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L4 AND PESTICIDE	44

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Derwent World Patents Index
IBM Technical Disclosure Bulletins

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L6

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DB=PGPB,USPT,USOC,EPAB,JPAB,DWPI,TDBD; PLUR=YES; OP=ADJ

<u>L6</u>	L4 AND PESTICIDE	44	<u>L6</u>
<u>L5</u>	L4 AND PESTCIDE	0	<u>L5</u>
<u>L4</u>	MALONONITRIL\$7.TI.	328	<u>L4</u>
<u>L3</u>	MALONONITRIL\$7	4001	<u>L3</u>

DB=PGPB; PLUR=YES; OP=ADJ

<u>L2</u>	20060004092	1	<u>L2</u>
<u>L1</u>	2006004092	0	<u>L1</u>

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☐ 1. Document ID: US 20070117854 A1

L6: Entry 1 of 44

File: PGPB

May 24, 2007

PGPUB-DOCUMENT-NUMBER: 20070117854

PGPUB-FILING-TYPE:

DOCUMENT-IDENTIFIER: US 20070117854 A1

TITLE: Malononitrile compounds and use thereof

PUBLICATION-DATE: May 24, 2007

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY
Mitsudera; Hiromasa	Toyonaka-shi		JP

US-CL-CURRENT: 514/383; 548/267.4

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	KWIC	Draw D
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☐ 2. Document ID: US 20060004092 A1

L6: Entry 2 of 44

File: PGPB

Jan 5, 2006

PGPUB-DOCUMENT-NUMBER: 20060004092

PGPUB-FILING-TYPE:

DOCUMENT-IDENTIFIER: US 20060004092 A1

TITLE: Malononitrile compound and use thereof pesticides

PUBLICATION-DATE: January 5, 2006

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY
Okada; Satoshi	Takarazuka-shi		JP
Oohra; Daisuke	Toyonaka-shi		JP
Otaka; Ken	Iwaki-shi		JP

US-CL-CURRENT: 514/521; 558/441

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	KWIC	Draw D
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☐ 3. Document ID: US 20050209323 A1

L6: Entry 3 of 44

File: PGPB

Sep 22, 2005

PGPUB-DOCUMENT-NUMBER: 20050209323

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20050209323 A1

TITLE: Malononitrile compounds and their use as pesticides

PUBLICATION-DATE: September 22, 2005

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY
Otaka, Ken	Osaka		JP
Oohira, Daisuke	Osaka		JP
Okada, Satoshi	Takarazuka-shi		JP

US-CL-CURRENT: 514/520; 558/388

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	KMOC	Draw D
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☐ 4. Document ID: US 20050176784 A1

L6: Entry 4 of 44

File: PGPB

Aug 11, 2005

PGPUB-DOCUMENT-NUMBER: 20050176784

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20050176784 A1

TITLE: Malononitrile compounds and their use as pesticides

PUBLICATION-DATE: August 11, 2005

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY
Otaka, Ken	Iwaki-shi		JP
Oohira, Daisuke	Toyonaka-shi		JP
Takaoka, Daisuke	Toyonaka-shi		JP

US-CL-CURRENT: 514/357; 546/320

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	KMOC	Draw D
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☐ 5. Document ID: US 20040143007 A1

L6: Entry 5 of 44

File: PGPB

Jul 22, 2004

PGPUB-DOCUMENT-NUMBER: 20040143007

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20040143007 A1

TITLE: Malononitrile compounds and their use as pesticides

PUBLICATION-DATE: July 22, 2004

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY
Otaka, Ken	Toyonaka-shi		JP
Oohira, Daisuke	Minoo-shi		JP
Suzuki, Masaya	Nishitokyo-shi		JP

US-CL-CURRENT: 514/520; 558/409

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	KMC	Draw D
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6. Document ID: US 20040142821 A1

L6: Entry 6 of 44

File: PGPB

Jul 22, 2004

PGPUB-DOCUMENT-NUMBER: 20040142821

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20040142821 A1

TITLE: Pesticide composition comprising malononitrile compounds

PUBLICATION-DATE: July 22, 2004

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY
Otaka, Ken	Toyonaka-shi		JP
Suzuki, Masaya	Nishitokyo-shi		JP
Oohira, Daisuke	Minoo-shi		JP

US-CL-CURRENT: 504/296; 504/309

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	KMC	Draw D
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7. Document ID: US 20040138065 A1

L6: Entry 7 of 44

File: PGPB

Jul 15, 2004

PGPUB-DOCUMENT-NUMBER: 20040138065

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20040138065 A1

TITLE: Malononitrile compounds and their use as pesticides

PUBLICATION-DATE: July 15, 2004

INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY
Otaka, Ken	Osaka		JP
Oohira, Daisuke	Osaka		JP
Okada, Satoshi	Hyogo		JP

US-CL-CURRENT: 504/309; 558/388

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	KMOC	Draw D
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☐ 8. Document ID: US 7026340 B2

L6: Entry 8 of 44

File: USPT

Apr 11, 2006

US-PAT-NO: 7026340

DOCUMENT-IDENTIFIER: US 7026340 B2

TITLE: Malononitrile compounds and their use as pesticides

PRIOR-PUBLICATION:

DOC-ID

DATE

US 20050176784 A1

August 11, 2005

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	KMOC	Draw D
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☐ 9. Document ID: US 7011838 B2

L6: Entry 9 of 44

File: USPT

Mar 14, 2006

US-PAT-NO: 7011838

DOCUMENT-IDENTIFIER: US 7011838 B2

TITLE: Malononitrile compounds and their use as pesticides

PRIOR-PUBLICATION:

DOC-ID

DATE

US 20040138065 A1

July 15, 2004

Full	Title	Citation	Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	KMOC	Draw D
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☐ 10. Document ID: US 6353126 B1

L6: Entry 10 of 44

File: USPT

Mar 5, 2002

US-PAT-NO: 6353126

DOCUMENT-IDENTIFIER: US 6353126 B1

TITLE: Process for the production of malononitrile

Full	Title	Citation	Front	Review	Classification	Date	Reference	Full Text	Abstract	Claims	KIMC	Draw D
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☐ 11. Document ID: US 6297393 B1

L6: Entry 11 of 44

File: USPT

Oct 2, 2001

US-PAT-NO: 6297393

DOCUMENT-IDENTIFIER: US 6297393 B1

TITLE: Process for the preparation of malononitrile

Full	Title	Citation	Front	Review	Classification	Date	Reference			Claims	KWMC	Draw D
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☐ 12. Document ID: WO 2006122949 A1

L6: Entry 12 of 44

File: EPAB

Nov 23, 2006

PUB-NO: WO2006122949A1

DOCUMENT-IDENTIFIER: WO 2006122949 A1

TITLE: MALONONITRILES AND THEIR USE AS PESTICIDES

Full	Title	Citation	Front	Review	Classification	Date	Reference			Claims	KWMC	Draw D
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☐ 13. Document ID: EP 1710234 A1

L6: Entry 13 of 44

File: EPAB

Oct 11, 2006

PUB-NO: EP001710234A1

DOCUMENT-IDENTIFIER: EP 1710234 A1

TITLE: MALONONITRILE COMPOUND AND USE THEREOF

Full	Title	Citation	Front	Review	Classification	Date	Reference			Claims	KWMC	Draw D
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☐ 14. Document ID: EP 1704143 A1

L6: Entry 14 of 44

File: EPAB

Sep 27, 2006

PUB-NO: EP001704143A1

DOCUMENT-IDENTIFIER: EP 1704143 A1

TITLE: MALONONITRILE COMPOUND AS PESTICIDES

Full	Title	Citation	Front	Review	Classification	Date	Reference			Claims	KWMC	Draw D
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☐ 15. Document ID: WO 2005068432 A1

L6: Entry 15 of 44

File: EPAB

Jul 28, 2005

PUB-NO: WO2005068432A1

DOCUMENT-IDENTIFIER: WO 2005068432 A1

TITLE: MALONONITRILE COMPOUND AS PESTICIDES

Full	Title	Citation	Front	Review	Classification	Date	Reference	Claims	KMOC	Draw D
------	-------	----------	-------	--------	----------------	------	-----------	--------	------	--------

☐ 16. Document ID: EP 1555259 A1

L6: Entry 16 of 44

File: EPAB

Jul 20, 2005

PUB-NO: EP001555259A1

DOCUMENT-IDENTIFIER: EP 1555259 A1

TITLE: Malononitrile compounds as pesticides

Full	Title	Citation	Front	Review	Classification	Date	Reference	Claims	KMOC	Draw D
------	-------	----------	-------	--------	----------------	------	-----------	--------	------	--------

☐ 17. Document ID: WO 2004020399 A1

L6: Entry 17 of 44

File: EPAB

Mar 11, 2004

PUB-NO: WO2004020399A1

DOCUMENT-IDENTIFIER: WO 2004020399 A1

TITLE: MALONONITRILE COMPOUND AND USE THEREOF AS PESTICIDES

Full	Title	Citation	Front	Review	Classification	Date	Reference	Claims	KMOC	Draw D
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☐ 18. Document ID: WO 2004006677 A1

L6: Entry 18 of 44

File: EPAB

Jan 22, 2004

PUB-NO: WO2004006677A1

DOCUMENT-IDENTIFIER: WO 2004006677 A1

TITLE: MALONONITRILE COMPOUNDS AND THEIR USE AS PESTICIDES

Full	Title	Citation	Front	Review	Classification	Date	Reference	Claims	KMOC	Draw D
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☐ 19. Document ID: WO 2089579 A1

L6: Entry 19 of 44

File: EPAB

Nov 14, 2002

PUB-NO: WO002089579A1

DOCUMENT-IDENTIFIER: WO 2089579 A1

TITLE: PESTICIDE COMPOSITION COMPRISING MALONONITRILE COMPOUNDS

Full	Title	Citation	Front	Review	Classification	Date	Reference	Claims	KMOC	Draw D
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☐ 20. Document ID: JP 2007055901 A

L6: Entry 20 of 44

File: DWPI

Mar 8, 2007

DERWENT-ACC-NO: 2007-317065

DERWENT-WEEK: 200731

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TITLE: Agent useful for preventing pest on livestock, comprises malononitrile compound and metoxadiazone as active ingredients

Full	Title	Citation	Front	Review	Classification	Date	Reference			Claims	KM/C	Draw D
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☐ 21. Document ID: JP 2007031422 A

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File: DWPI

Feb 8, 2007

DERWENT-ACC-NO: 2007-246448

DERWENT-WEEK: 200725

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TITLE: New malononitrile compound useful for controlling pests, insects, mites and nematodes

Full	Title	Citation	Front	Review	Classification	Date	Reference	Claims	KMOC	Draw D
------	-------	----------	-------	--------	----------------	------	-----------	--------	------	--------

☐ 22. Document ID: JP 2007001934 A

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File: DWPI

Jan 11, 2007

DERWENT-ACC-NO: 2007-211549

DERWENT-WEEK: 200722

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TITLE: Composition for controlling pests such as Blattodea insect-pest and flies, contains malononitrile compound and natural pyrethrin as active ingredients

Full	Title	Citation	Front	Review	Classification	Date	Reference	Claims	KMOC	Draw D
------	-------	----------	-------	--------	----------------	------	-----------	--------	------	--------

☐ 23. Document ID: JP 2007001935 A

L6: Entry 23 of 44

File: DWPI

Jan 11, 2007

DERWENT-ACC-NO: 2007-165230

DERWENT-WEEK: 200717

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TITLE: Pest controller for controlling harmful pests such as Blattodea insect-pest and flies in agriculture and forestry, contains malononitrile compound and ethofenprox as active ingredients

Full	Title	Citation	Front	Review	Classification	Date	Reference	Claims	KMOC	Draw D
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☐ 24. Document ID: JP 2007001933 A

L6: Entry 24 of 44

File: DWPI

Jan 11, 2007

DERWENT-ACC-NO: 2007-165229

DERWENT-WEEK: 200717

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TITLE: Pest controller for controlling harmful pests such as Blattodea insect-pest and flies in agriculture and forestry, contains malononitrile compound and bifenthrin as active ingredients

Full	Title	Citation	Front	Review	Classification	Date	Reference	Abstract	Summary	Claims	KWIC	Draw D
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☐ 25. Document ID: JP 2007001932 A

L6: Entry 25 of 44

File: DWPI

Jan 11, 2007

DERWENT-ACC-NO: 2007-154719

DERWENT-WEEK: 200716

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TITLE: Composition for controlling pests such as Blattodea insect-pest and flies, contains malononitrile compound and empenhrin as active ingredients

Full	Title	Citation	Front	Review	Classification	Date	Reference	Abstract	Summary	Claims	KWIC	Draw D
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☐ 26. Document ID: JP 2007001931 A

L6: Entry 26 of 44

File: DWPI

Jan 11, 2007

DERWENT-ACC-NO: 2007-154718

DERWENT-WEEK: 200716

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TITLE: Composition for controlling pests such as Blattodea insect-pest and flies, contains malononitrile compound and imiprothrin as active ingredients

Full	Title	Citation	Front	Review	Classification	Date	Reference	Abstract	Summary	Claims	KWIC	Draw D
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☐ 27. Document ID: JP 2007001930 A

L6: Entry 27 of 44

File: DWPI

Jan 11, 2007

DERWENT-ACC-NO: 2007-154717

DERWENT-WEEK: 200716

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TITLE: Composition for controlling pests such as Blattodea insect-pest and flies, contains malononitrile compound and allethrin or prallethrin as active ingredients

Full	Title	Citation	Front	Review	Classification	Date	Reference	Abstract	Summary	Claims	KWIC	Draw D
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☐ 28. Document ID: JP 2007001929 A

L6: Entry 28 of 44

File: DWPI

Jan 11, 2007

DERWENT-ACC-NO: 2007-154716

DERWENT-WEEK: 200716

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TITLE: Composition for controlling pests such as flies, ticks and moths, contains malononitrile compound and 2,3,5,6-tetrafluoro benzyl ester compound as active ingredients

Full	Title	Citation	Front	Review	Classification	Date	Reference	Abstract	Summary	Claims	KWIC	Draw D
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☐ 29. Document ID: JP 2007001928 A

L6: Entry 29 of 44

File: DWPI

Jan 11, 2007

DERWENT-ACC-NO: 2007-135315

DERWENT-WEEK: 200714

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TITLE: Composition for controlling pests such as Blattodea insect-pest, flies, ticks and moths, contains malononitrile compound and 3-phenoxy benzyl ester compound as active ingredients

Full	Title	Citation	Front	Review	Classification	Date	Reference	Abstract	Summary	Claims	KWIC	Draw D
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☐ 30. Document ID: CN 1910156 A, WO 2005068432 A1, JP 2005225875 A, EP 1704143 A1, BR 200506447 A, AU 2005205311 A1

L6: Entry 30 of 44

File: DWPI

Feb 7, 2007

DERWENT-ACC-NO: 2005-571136

DERWENT-WEEK: 200743

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TITLE: New malononitrile compounds used e.g. in pesticidal composition for controlling pest and for controlling parasites living outside of livestock

Full	Title	Citation	Front	Review	Classification	Date	Reference	Abstract	Summary	Claims	KWIC	Draw D
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☐ 31. Document ID: CN 1910147 A, WO 2005068423 A1, JP 2005225876 A, EP 1710234 A1, AU 2005205298 A1, BR 200506451 A, US 20070117854 A1, KR 2006127968 A

L6: Entry 31 of 44

File: DWPI

Feb 7, 2007

DERWENT-ACC-NO: 2005-563927

DERWENT-WEEK: 200743

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TITLE: New malononitrile compound for preventing pest such as Hemiptera insect-pest, planthoppers, leafhoppers, plant louses, stink bugs, greenhouse whitefly and scale insects

Full	Title	Citation	Front	Review	Classification	Date	Reference			Claims	KWIC	Draw. De
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☐ 32. Document ID: EP 1555259 A1, JP 2005225873 A, CN 1648122 A, =I N 20040150 0I

L6: Entry 32 of 44

File: DWPI

Jul 20, 2005

DERWENT-ACC-NO: 2005-513905

DERWENT-WEEK: 200641

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TITLE: New malononitrile derivatives, useful in pesticide composition for controlling pests living external to livestock such as cow or horse

Full	Title	Citation	Front	Review	Classification	Date	Reference			Claims	KWIC	Draw. De
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☐ 33. Document ID: JP 2004099597 A

L6: Entry 33 of 44

File: DWPI

Apr 2, 2004

DERWENT-ACC-NO: 2004-299646

DERWENT-WEEK: 200428

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TITLE: Pest controller for controlling pest e.g. harmful arthropods, insects, mites and nematodes, and for preventing ectoparasite in animals e.g. cow, pig, sheep and dog, contains malononitrile derivatives as active ingredient

Full	Title	Citation	Front	Review	Classification	Date	Reference			Claims	KWIC	Draw. De
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☐ 34. Document ID: JP 2004099593 A

L6: Entry 34 of 44

File: DWPI

Apr 2, 2004

DERWENT-ACC-NO: 2004-299644

DERWENT-WEEK: 200428

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TITLE: New malononitrile compounds useful in exterminating pests, arthropods, insects, mites, nematodes and parasites, is new

Full	Title	Citation	Front	Review	Classification	Date	Reference	Abstract	Summary	Claims	KM/C	Draw D
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☐ 35. Document ID: ES 2276090 T3, WO 2004006677 A1, JP 2004099592 A, AU 2003281174 A1, EP 1521528 A1, BR 200312638 A, KR 2005019888 A, US 20050176784 A1, CN 1668195 A, IN 200403066 P4, US 7026340 B2, EP 1521528 B1, DE 60309874 E, DE 60309874 T2

L6: Entry 35 of 44

File: DWPI

Jun 16, 2007

DERWENT-ACC-NO: 2004-191068

DERWENT-WEEK: 200742

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TITLE: New malononitrile compounds useful for controlling pests e.g. insect pests, acarine pests and nematode pests in agriculture and forestry

Full	Title	Citation	Front	Review	Classification	Date	Reference	Abstract	Summary	Claims	KM/C	Draw D
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☐ 36. Document ID: AU 2002307747 A1, WO 200290321 A1, EP 1392647 A1, BR 200209481 A, US 20040143007 A1

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File: DWPI

Nov 18, 2002

DERWENT-ACC-NO: 2003-093276

DERWENT-WEEK: 200452

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TITLE: New malononitrile compounds useful for controlling pests e.g. insect pests

Full	Title	Citation	Front	Review	Classification	Date	Reference	Abstract	Summary	Claims	KM/C	Draw D
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☐ 37. Document ID: AU 2002255313 B2, WO 200290320 A2, EP 1385817 A2, BR 200209532 A, HU 200400033 A2, US 20040138065 A1, AU 2002255313 A1, US 7011838 B2, RU 2274638 C2

L6: Entry 37 of 44

File: DWPI

Feb 1, 2007

DERWENT-ACC-NO: 2003-093275

DERWENT-WEEK: 200735

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TITLE: New malononitrile compounds useful for controlling pests e.g. insect pests

Full	Title	Citation	Front	Review	Classification	Date	Reference	Claims	KWIC	Draw. De
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☐ 38. Document ID: IN 200301743 P4, WO 200289579 A1, JP 2003026510 A, JP 2003026511 A, JP 2003026647 A, EP 1385377 A1, KR 2003092137 A, KR 2003092138 A, KR 2004012807 A, BR 200209461 A, US 20040142821 A1, AU 2002307746 A1, CN 1523958 A, CN 1524071 A, TW 223979 B1, US 20050209323 A1, CN 1639114 A

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File: DWPI

Jan 6, 2006

DERWENT-ACC-NO: 2003-093225

DERWENT-WEEK: 200615

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TITLE: Composition useful for controlling pests comprises a malononitrile compound

Full	Title	Citation	Front	Review	Classification	Date	Reference	Claims	KWIC	Draw. De
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☐ 39. Document ID: DE 60012918 T2, US 6297393 B1, EP 1184369 A1, JP 2002145843 A, EP 1184369 B1, DE 60012918 E

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File: DWPI

Sep 8, 2005

DERWENT-ACC-NO: 2002-040115

DERWENT-WEEK: 200559

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TITLE: Preparation of malononitrile from cyanoacetamide useful as building block for pharmaceuticals and pesticides, e.g. thiamine, adenine comprises porous particulate solid substance as absorbent replacing inorganic salt.

Full	Title	Citation	Front	Review	Classification	Date	Reference	Claims	KWIC	Draw. De
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☐ 40. Document ID: DE 60005252 E, EP 1065198 A1, CA 2312514 A1, JP 2001039939 A, CN 1296944 A, US 6353126 B1, EP 1065198 B1

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File: DWPI

Oct 23, 2003

DERWENT-ACC-NO: 2001-184252

DERWENT-WEEK: 200377

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TITLE: Preparation of malononitrile by reacting cyanoacetamide with cyanuric chloride in the presence of a catalytic amount of N,N-dimethylformamide in polar solvent

Full	Title	Citation	Front	Review	Classification	Date	Reference	Claims	KWIC	Draw. De
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Terms	Documents
L4 AND PESTICIDE	44

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Search Results - Record(s) 41 through 44 of 44 returned.

☐ 41. Document ID: CN 1035874 C, EP 611751 A1, US 5344955 A, AU 9455247 A, BR 9400605 A, CA 2115908 A, CZ 9400355 A3, JP 06256286 A, SK 9400189 A3, ZA 9401114 A, NZ 250918 A, AU 671334 B, EP 611751 B1, DE 69402092 E, ES 2100657 T3, CN 1107465 A, IL 108678 A, RU 2111960 C1, MX 186472 B, SG 49606 A1, SK 280168 B6, CZ 289315 B6, KR 285611 B, CA 2115908 C, JP 3512844 B2

L6: Entry 41 of 44

File: DWPI

Sep 17, 1997

DERWENT-ACC-NO: 1994-265362

DERWENT-WEEK: 200455

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TITLE: Prepn. of sodium salt of 1-amino-1-cyanamido-2,2- di:cyano ethylene - from malononitrile and sodium di:cyanamide in aprotic, dipolar solvent, at elevated temp.

Full	Title	Citation	Front	Review	Classification	Date	Reference	Claims	KWIC	Draw D
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☐ 42. Document ID: EP 508353 A1, DE 59206635 G, CA 2065765 A, JP 05112535 A, US 5216160 A, EP 508353 B1

L6: Entry 42 of 44

File: DWPI

Oct 14, 1992

DERWENT-ACC-NO: 1992-341662

DERWENT-WEEK: 199636

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TITLE: 4-Amino-2-chloro-5-cyano-6-methylthio-pyrimidine prodn. - by reacting malononitrile and carbon di:sulphide with strong base, further reacting with methylating agent condensing with cyanamide and cyclising with hydrochloric acid

Full	Title	Citation	Front	Review	Classification	Date	Reference	Claims	KWIC	Draw D
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☐ 43. Document ID: DE 2601052 A, BE 837583 A, NL 7600379 A, JP 51095029 A, FR 2297838 A, US 4000314 A, ZA 7600227 A, AT 7600244 A, GB 1475974 A, IL 48847 A, CH 604508 A, CA 1061799 A, SU 708979 A

L6: Entry 43 of 44

File: DWPI

Jul 22, 1976

DERWENT-ACC-NO: 1976-58134X

DERWENT-WEEK: 200300

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TITLE: ((N-Benzyl-3-trifluoromethyl-anilino)methylene)malonitrile derivs - prepd by reacting trifluoromethylanilinomethylene-malononitriles with benzyl halides (BE150776)

Full	Title	Citation	Front	Review	Classification	Date	Reference	Claims	KMOC	Draw D
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☐ 44. Document ID: US 3726662 A

L6: Entry 44 of 44

File: USOC

Apr 10, 1973

US-PAT-NO: 3726662

DOCUMENT-IDENTIFIER: US 3726662 A

TITLE: HERBICIDAL ANILINOMETHYLENE-MALONONITRILES

DATE-ISSUED: April 10, 1973

INVENTOR-NAME: BAKER J; HOWE R

US-CL-CURRENT: 504/312

Full	Title	Citation	Front	Review	Classification	Date	Reference	Claims	KMOC	Draw D
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L4 AND PESTICIDE

44

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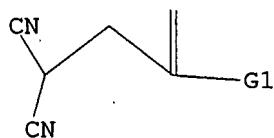
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FILE LAST UPDATED: 30 Jul 2007 (20070730/ED)

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<http://www.cas.org/infopolicy.html>

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L1 STR



G1 O,S

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=> s 11
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Substance data SEARCH and crossover from CAS REGISTRY in progress...
Use DISPLAY HITSTR (or FHITSTR) to directly view retrieved structures.

SAMPLE SEARCH INITIATED 15:33:52 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED - 63 TO ITERATE

100.0% PROCESSED 63 ITERATIONS 13 ANSWERS
SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**
BATCH **COMPLETE**
PROJECTED ITERATIONS: 784 TO 1736
PROJECTED ANSWERS: 44 TO 476

L2 13 SEA SSS SAM L1

L3 10 L2

=> d 1-10 ibib abs hitstr

L3 ANSWER 1 OF 10 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 2004:203802 CAPLUS

DOCUMENT NUMBER: 140:235428

TITLE: Preparation of malononitrile compound and use thereof as pesticides

INVENTOR(S): Okada, Satoshi; Oohira, Daisuke; Otaka, Ken

PATENT ASSIGNEE(S): Sumitomo Chemical Company, Limited, Japan

SOURCE: PCT Int. Appl., 104 pp.

CODEN: PIXXD2

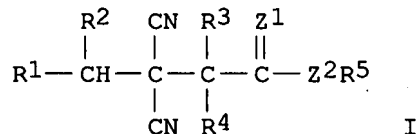
DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

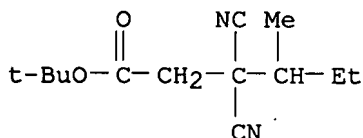
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004020399	A1	20040311	WO 2003-JP10726	20030826
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, KE, KG, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW			
RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			
AU 2003256083	A1	20040319	AU 2003-256083	20030826
BR 2003013964	A	20050719	BR 2003-13964	20030826
CN 1678571	A	20051005	CN 2003-820424	20030826
JP 2004143148	A	20040520	JP 2003-208994	20030827
US 2006004092	A1	20060105	US 2005-522764	20050201
PRIORITY APPLN. INFO.:			JP 2002-250355	A 20020829
			WO 2003-JP10726	W 20030826
OTHER SOURCE(S):	MARPAT 140:235428			
GI				



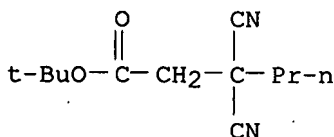
AB The present invention relates to a novel malononitrile compound represented by the formula (I): wherein, R1 represents C1 to C6 alkyl that may be substituted with halogen, C2 to C6 alkenyl that may be substituted with halogen, etc; R2 represents hydrogen atom or C1 to C6 alkyl that may be substituted with halogen; R3 represents hydrogen atom or C1 to C6 alkyl; R4 represents hydrogen atom or C1 to C6 alkyl; R5 represents C1 to C6 alkyl that may be substituted with halogen, C3 to C6 alkenyl that may be substituted with halogen, etc, or R4 and R5 may be combined at their terminal and represent ethylene that may be substituted with C1 to C3 alkyl or trimethylene that may be substituted with C1 to C3 alkyl; and Z1 and Z2, which are the same or different, represent oxygen atom or sulfur atom. Thus, 2-(tert-butoxycarbonylmethyl)-2-allylmalononitrile was prepared by reacting 2-allylmalonitrile with tert-Bu bromoacetate in DMF in the

presence of sodium hydride. The malononitrile compound has an efficient pesticidal activity and can control effectively pests such as insect pests, acarine pests, nematode pests and the like.

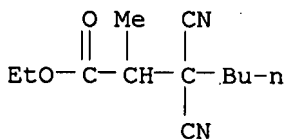
- IT 666738-88-1P, 2-(tert-Butoxycarbonylmethyl)-2-(1-methylpropyl)malononitrile 666738-93-8P, 2-(tert-Butoxycarbonylmethyl)-2-propylmalononitrile 666738-94-9P, 2-[1-(Ethoxycarbonyl)ethyl]-2-butylmalononitrile 666738-97-2P 666739-14-6P, 2-[(3-Methyl-3-methoxybutoxy)carbonylmethyl]-2-(3,3,3-trifluoropropyl)malononitrile
 RL: AGR (Agricultural use); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses)
 (production of malononitriles as pesticides)
 RN 666738-88-1 CAPLUS
 CN Hexanoic acid, 3,3-dicyano-4-methyl-, 1,1-dimethylethyl ester (9CI) (CA INDEX NAME)



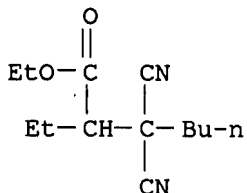
- RN 666738-93-8 CAPLUS
 CN Hexanoic acid, 3,3-dicyano-, 1,1-dimethylethyl ester (9CI) (CA INDEX NAME)



- RN 666738-94-9 CAPLUS
 CN Heptanoic acid, 3,3-dicyano-2-methyl-, ethyl ester (9CI) (CA INDEX NAME)

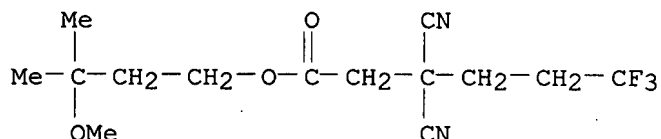


- RN 666738-97-2 CAPLUS
 CN Heptanoic acid, 3,3-dicyano-2-ethyl-, ethyl ester (9CI) (CA INDEX NAME)



- RN 666739-14-6 CAPLUS
 CN Hexanoic acid, 3,3-dicyano-6,6,6-trifluoro-, 3-methoxy-3-methylbutyl ester

(9CI) (CA INDEX NAME)



REFERENCE COUNT: 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 2 OF 10 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 2000:141061 CAPLUS

DOCUMENT NUMBER: 132:278722

TITLE: Spontaneous addition of active methine compounds to enol ethers and α,β -unsaturated ketones in aprotic polar solvent

AUTHOR(S): Yokozawa, Tsutomu; Oishi, Motoi; Tanaka, Yasukazu

CORPORATE SOURCE: Department of Applied Chemistry, Kanagawa University, Kanagawa-ku Yokohama, 221-8686, Japan

SOURCE: Journal of Organic Chemistry (2000), 65(6), 1895-1897
CODEN: JOCEAH; ISSN: 0022-3263

PUBLISHER: American Chemical Society

DOCUMENT TYPE: Journal

LANGUAGE: English

OTHER SOURCE(S): CASREACT 132:278722.

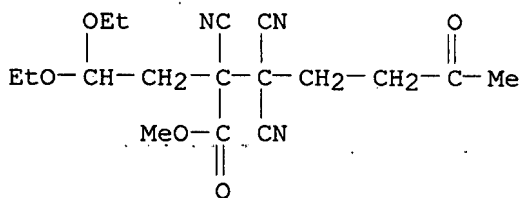
AB Addition of (EtO)2CHCH2CX(Y)CH(CN)2 (I, X = Y = cyano; X = cyano, Y = CO2Me; X = Y = CO2Me) to enol ethers and α,β -unsatd. ketones in DMF at room temp is reported. Thus, reacting I (X = Y = cyano) with H2C:CHOEt gave (EtO)2CHCH2C(CN)2CH(OEt)Me in 63% yield. This reaction illustrates that the electron-withdrawing groups at the β -positions of the active methine group having the ones at the α and β positions were strongly affected on the acidity of I.

IT 264142-40-7P 264142-41-8P 264142-43-0P

RL: SPN (Synthetic preparation); PREP (Preparation)
(addition of methine compds. to enol ethers and α,β -unsatd. ketones)

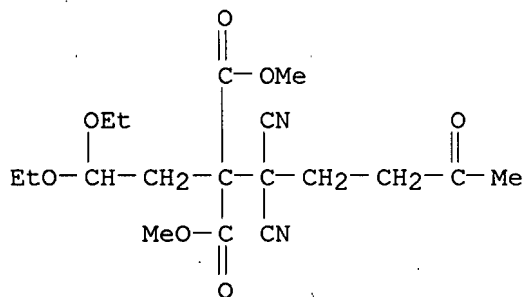
RN 264142-40-7 CAPLUS

CN Heptanoic acid, 2,3,3-tricyano-2-(2,2-diethoxyethyl)-6-oxo-, methyl ester (9CI) (CA INDEX NAME)

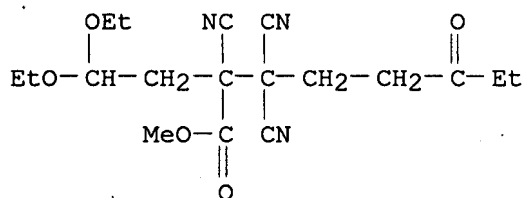


RN 264142-41-8 CAPLUS

CN Propanedioic acid, (1,1-dicyano-4-oxopentyl)(2,2-diethoxyethyl)-, dimethyl ester (9CI) (CA INDEX NAME)



RN 264142-43-0 CAPLUS
 CN Octanoic acid, 2,3,3-tricyano-2-(2,2-diethoxyethyl)-6-oxo-, methyl ester
 (9CI) (CA INDEX NAME)



REFERENCE COUNT: 25 THERE ARE 25 CITED REFERENCES AVAILABLE FOR THIS
 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 3 OF 10 CAPLUS COPYRIGHT 2007 ACS on STN
 ACCESSION NUMBER: 1999:142376 CAPLUS
 DOCUMENT NUMBER: 130:239567
 TITLE: Diazaspirononanium salt for use as template for
 zeolite synthesis
 INVENTOR(S): Kubota, Yoshihiro; Sugi, Yoshihiro
 PATENT ASSIGNEE(S): Showa Denko K. K., Japan
 SOURCE: Jpn. Kokai Tokkyo Koho, 9 pp.
 CODEN: JKXXAF
 DOCUMENT TYPE: Patent
 LANGUAGE: Japanese
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

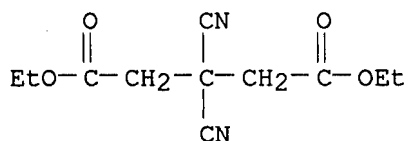
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 11060577	A	19990302	JP 1997-220414	19970815
PRIORITY APPLN. INFO.:			JP 1997-220414	19970815

OTHER SOURCE(S): MARPAT 130:239567

AB Claimed template is a salt of substituted 2,7-diazaspiro[4,4]nonanium.
 Hydrothermal synthesis of a zeolite by bringing a silica source and/or an
 alumina source into contact with the zeolite is also claimed. ZSM-12
 zeolites having crystal size of a major axis $\geq 50 \mu\text{m}$ are also
 claimed.

IT 77415-69-1P
 RL: IMF (Industrial manufacture); RCT (Reactant); PREP (Preparation); RACT
 (Reactant or reagent)
 (reaction of; diazaspirononanium salts as templates for manufacture of
 ZSM-12 zeolites having large crystal size)

RN 77415-69-1 CAPLUS
 CN Pentanedioic acid, 3,3-dicyano-, diethyl ester (9CI) (CA INDEX NAME)



L3 ANSWER 4 OF 10 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1995:900662 CAPLUS

DOCUMENT NUMBER: 124:116317

TITLE: Lanthanum isopropoxide catalyzed addition of activated nucleophiles to imines

AUTHOR(S): Yamamoto, Yoshinori; Fukui, Hiroyuki; Honda, Yoshihiro

CORPORATE SOURCE: Dept. Chem., Tohoku Univ., Sendai, 980-77, Japan

SOURCE: Applied Organometallic Chemistry (1995), 9(5 & 6), 467-71

CODEN: AOCHEX; ISSN: 0268-2605

PUBLISHER: Wiley

DOCUMENT TYPE: Journal

LANGUAGE: English

OTHER SOURCE(S): CASREACT 124:116317

AB The addition of certain activated nucleophiles to activated imines is catalyzed by lanthanum isopropoxide. As activated nucleophiles, methylmalonitrile and Me 2-cyanopropanoate can be utilized. Imines having an electron-withdrawing group either at the carbon or at the nitrogen atom of the C:N double bond can be used: for example N-toluenesulfonylimines, N-(4-methoxycarbonylphenyl)imines and α -imino esters.

IT 173006-25-2P

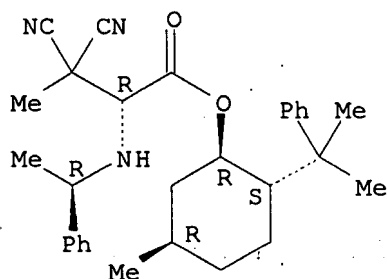
RL: SPN (Synthetic preparation); PREP (Preparation)

(lanthanum isopropoxide catalyzed addition of activated nucleophiles to imines)

RN 173006-25-2 CAPLUS

CN Butanoic acid, 3,3-dicyano-2-[(1-phenylethyl)amino]-, 5-methyl-2-(1-methyl-1-phenylethyl)cyclohexyl ester, [1R-[1 α [R*(R*)],2 β ,5 α]]-(9CI) (CA INDEX NAME)

Absolute stereochemistry.




L3 ANSWER 5 OF 10 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1992:83465 CAPLUS

DOCUMENT NUMBER: 116:83465

TITLE: The regioselectivity of the ring opening of 1-activated or nonactivated 2-alkoxycarbonyl or 2-cyanoaziridines by carbanions of the dicarbonyl compounds

AUTHOR(S): Bouayad, Zoheir; Chanet-Ray, Josette; Ducher, S.; Vessiere, Roger

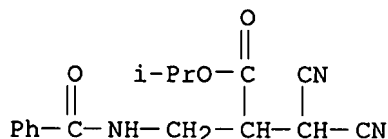


AB Aziridines, e.g. I, reacted with carbanions of dicarbonyl compds., e.g. $\text{RO}_2\text{CCH}_2\text{CO}_2\text{R}$ ($\text{R} = \text{Me}, \text{Et}, \text{CHMe}_2$), to give ring opened products and/or ring enlarged products, e.g. $(\text{RO}_2\text{C})_2\text{CHCH}_2\text{CH}(\text{NHBz})\text{CO}_2\text{CHMe}_2$, $(\text{RO}_2\text{C})_2\text{CHCH}(\text{CO}_2\text{CHMe}_2)\text{CH}_2\text{NHBz}$, and pyrrole II. The regioselectivity depends on several factors. The Ph group on C-3 favors C-3-N bond cleavage, whereas C-2-N bond cleavage is predominant with C-3 substituted or C-2-H aziridines. Cyanoaziridines are predominantly cleaved at C-3-N.

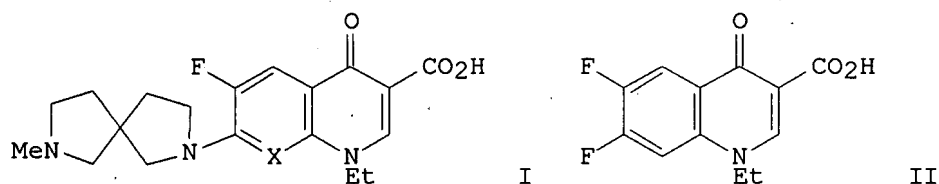
IT 138478-35-0P
RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of)

RN 138478-35-0 CAPLUS

CN Propanoic acid, 2-[(benzoylamino)methyl]-3,3-dicyano-, 1-methylethyl ester (9CI) (CA INDEX NAME)



L3 ANSWER 6 OF 10 CAPLUS COPYRIGHT 2007 ACS on STN
ACCESSION NUMBER: 1990:497432 CAPLUS
DOCUMENT NUMBER: 113:97432
TITLE: Quinolone antibacterial agents substituted at the
7-position with spiroamines. Synthesis and
structure-activity relationships
AUTHOR(S): Culbertson, Townley P.; Sanchez, Joseph P.; Gambino,
Laura; Sesnie, Josephine A.
CORPORATE SOURCE: Parke-Davis Pharm. Res. Div., Warner-Lambert Co., Ann Arbor, MI, 48105, USA
SOURCE: Journal of Medicinal Chemistry (1990), 33(8), 2270-5
CODEN: JMCMAR; ISSN: 0022-2623
DOCUMENT TYPE: Journal
LANGUAGE: English
OTHER SOURCE(S): CASREACT 113:97432
GI



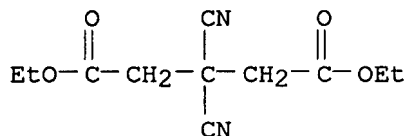
AB Fluoroquinolone antibacterials having the 7-position (10-position of pyridobenzoxazines) substituted with 2,7-diazaspiro[4.4]nonane, 1,7-diazaspiro[4.4]nonane, or 2,8-diazaspiro[5.5]undecane (e.g. I (X = CF, CH, N) were prepared and their biol. activities were compared with piperazine and pyrrolidine substituted analogs. Most exhibited potent Gram-pos. and Gram-neg. activity, especially when side chain was N-alkylated. Thus, the quinolinecarboxylic acid II was treated with 2-methyl-2,7-diazaspiro[4.4]nonane to give I (X = CH).

IT 77415-69-1

RL: RCT (Reactant); RACT (Reactant or reagent)
(reductive cyclization of)

RN 77415-69-1 CAPLUS

CN Pentanedioic acid, 3,3-dicyano-, diethyl ester (9CI) (CA INDEX NAME)



L3 ANSWER 7 OF 10 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1986:496793 CAPLUS

DOCUMENT NUMBER: 105:96793

TITLE: Zwitterionic tetramethylenes as the common intermediates in the cycloaddition and polymerization reactions of N-vinylcarbazole with electrophilic tetrasubstituted ethylenes: a new explanation for charge-transfer initiation

AUTHOR(S): Gotoh, Tetsuya; Padias, Anne Buyle; Hall, H. K., Jr.

CORPORATE SOURCE: Chem. Dep., Univ. Arizona, Tucson, AZ, 85721, USA

SOURCE: Journal of the American Chemical Society (1986), 108(16), 4920-31

CODEN: JACSAT; ISSN: 0002-7863

DOCUMENT TYPE: Journal

LANGUAGE: English

OTHER SOURCE(S): CASREACT 105:96793

AB The reactions of N-vinylcarbazole (I) with electrophilic tetrasubstituted ethylenes were examples of reactions whose outcomes are manipulated by changes in concentration, structure, and working procedure to form either small mols. (cyclobutanes, 1-butenes) or poly(vinylcarbazole). Equivalent concns. and evaporative workup (organic chemists' conditions) lead to small mols.; a large excess of I and precipitative workup give polymer. The mechanism involves gauche and trans zwitterionic tetramethylenes as intermediates. The former gives cyclobutane reversibly. The latter gives 1-butenes intramol. or adds monomers to form cyclohexanes or eventually polymer. The organic chemical and polymer chemical are unified on this basis. Extensive stereochem. and kinetic support for these propositions is given. Two other proposed mechanisms for these charge-transfer initiations are excluded.

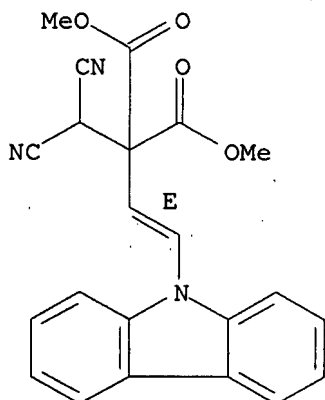
IT 96735-90-9P

RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of)

RN 96735-90-9 CAPLUS

CN Propanedioic acid, [2-(9H-carbazol-9-yl)ethenyl](dicyanomethyl)-, dimethyl ester, (E)- (9CI) (CA INDEX NAME)

Double bond geometry as shown.



L3 ANSWER 8 OF 10 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1985:406747 CAPLUS

DOCUMENT NUMBER: 103:6747

TITLE: Zwitterionic tetramethylene intermediates: a new interpretation for "charge-transfer" initiation

AUTHOR(S): Hall, H. K., Jr.; Gotoh, T.

CORPORATE SOURCE: Dep. Chem., Univ. Arizona, Tucson, AZ, 85721, USA

SOURCE: Polymer Preprints (American Chemical Society, Division of Polymer Chemistry) (1985), 26(1), 34-5
CODEN: ACPPAY; ISSN: 0032-3934

DOCUMENT TYPE: Journal

LANGUAGE: English

AB Investigation of the initiation mechanism in polymerization of N-vinylcarbazole (I) [1484-13-5] in the presence of tetracyanoethylene [670-54-2] or di-Me 2,2-dicyanoethylene-1,1-dicarboxylate [82849-49-8] showed that neither the I-cyano compound charge transfer complexes nor the ion-radical pairs formed from them initiated polymerization. The initiating species was the gauche or trans tetramethylene zwitterion formed as an intermediate from the charge-transfer complex. This finding indicated that cyclobutanes initiated vinyl polymerization. The mechanism and the kinetics of the zwitterionic initiation were discussed.

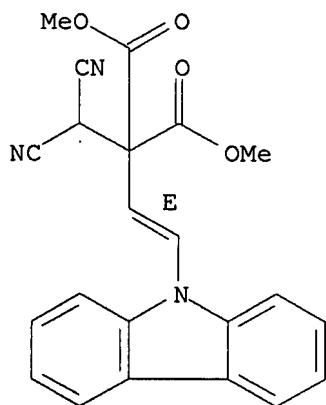
IT 96735-90-9

RL: CAT (Catalyst use); USES (Uses)
(catalysts, for vinylcarbazole polymerization)

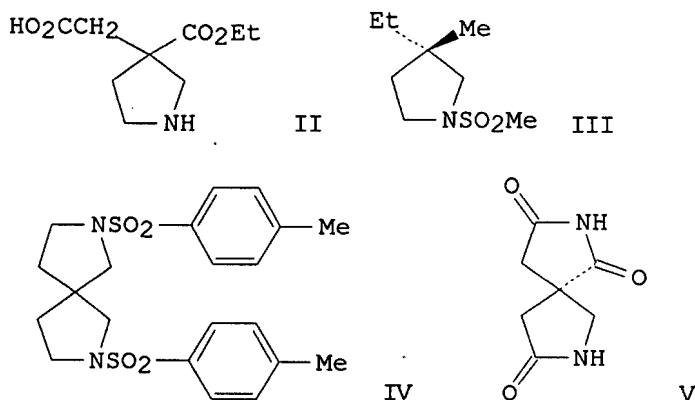
RN 96735-90-9 CAPLUS

CN Propanedioic acid, [2-(9H-carbazol-9-yl)ethenyl](dicyanomethyl)-, dimethyl ester, (E)- (9CI) (CA INDEX NAME)

Double bond geometry as shown.



L3 ANSWER 9 OF 10 CAPLUS COPYRIGHT 2007 ACS on STN
 ACCESSION NUMBER: 1981:442180 CAPLUS
 DOCUMENT NUMBER: 95:42180
 TITLE: Absolute configuration of 2,7-diazaspiro[4,4]nonane.
 A reassignment
 AUTHOR(S): Overberger, C. G.; Wang, David Wei; Hill, Richard K.;
 Krow, Grant R.; Ladner, David W.
 CORPORATE SOURCE: Macromol. Res. Cent., Univ. Michigan, Ann Arbor, MI,
 48109, USA
 SOURCE: Journal of Organic Chemistry (1981), 46(13), 2757-64
 CODEN: JOCEAH; ISSN: 0022-3263
 DOCUMENT TYPE: Journal
 LANGUAGE: English
 OTHER SOURCE(S): CASREACT 95:42180
 GI



AB The absolute configuration of the axially dissym. spirane 2,7-diazaspiro[4,4]nonane (I), was elucidated as (R)-(-), (S)-(+) in CHCl₃ by synthesis of both enantiomers from the centrodissym. intermediate II; the configuration of (R)-(-)-II was correlated with that of (S)-HO₂CCMeEtCH₂CO₂H through the substituted pyrrolidine III. The configuration thus established for the sulfonamide derivative IV is opposite to that derived earlier (Krow, G. and Hill, R. K., 1968). The source of the original error lies in the preparation of spiroimide V, which is accompanied by almost total racemization when carried out at high temps. A more direct, efficient synthesis of I is described, followed by resolution with dinitrodiphenic acid to give the optically pure enantiomers. Lowe's

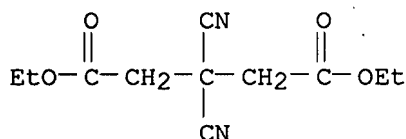
rule predicts correctly the absolute configurations of several I derivs. but not that of I itself.

IT 77415-69-1P

RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation and diazaspirononane derivative from)

RN 77415-69-1 CAPLUS

CN Pentanedioic acid, 3,3-dicyano-, diethyl ester (9CI) (CA INDEX NAME)



L3 ANSWER 10 OF 10 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1978:50444 CAPLUS

DOCUMENT NUMBER: 88:50444

TITLE: The chemistry of 2-oxopropanedinitrile (carbonyl cyanide); XIX. The ene synthesis using 2-oxopropanedinitrile and 1,3-dicarbonyl compounds
AUTHOR(S): Kociolek, K.; Leplawy, M. T.
CORPORATE SOURCE: Inst. Org. Chem., Tech. Univ. Lodz, Lodz, Pol.
SOURCE: Synthesis (1977), (11), 778-80
CODEN: SYNTBF; ISSN: 0039-7881

DOCUMENT TYPE: Journal

LANGUAGE: English

OTHER SOURCE(S): CASREACT 88:50444

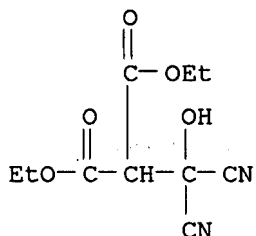
AB Reaction of $\text{CO}(\text{CN})_2$ with $\text{RCOCH}_2\text{COR}_1$ (I; $\text{R} = \text{R}_1 = \text{Ph}$, 2,4,6- $\text{Cl}_3\text{C}_6\text{H}_2$, Me; $\text{R} = \text{Me}$, F_3C , $\text{R}_1 = \text{Ph}$) in ether at 0° was complete in 1 h and gave $\text{RCOCH}(\text{COR}_1)\text{C}(\text{CN})_2\text{OH}$ (II; R and R_1 as before) in 100% yield. Reaction of $\text{CO}(\text{CN})_2$ with I ($\text{R} = \text{R}_1 = \text{OEt}$) at room temperature required 20 days and gave II in 43-66% yield.

IT 65305-78-4P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation and reaction of, with aniline)

RN 65305-78-4 CAPLUS

CN Propanedioic acid, (dicyanohydroxymethyl)-, diethyl ester (9CI) (CA INDEX NAME)



=> s 11 full

REGISTRY INITIATED

Substance data SEARCH and crossover from CAS REGISTRY in progress...

Use DISPLAY HITSTR (or FHITSTR) to directly view retrieved structures.

FULL SEARCH INITIATED 15:35:27 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 1125 TO ITERATE

100.0% PROCESSED 1125 ITERATIONS 208 ANSWERS
SEARCH TIME: 00.00.01

L4 208 SEA SSS FUL L1

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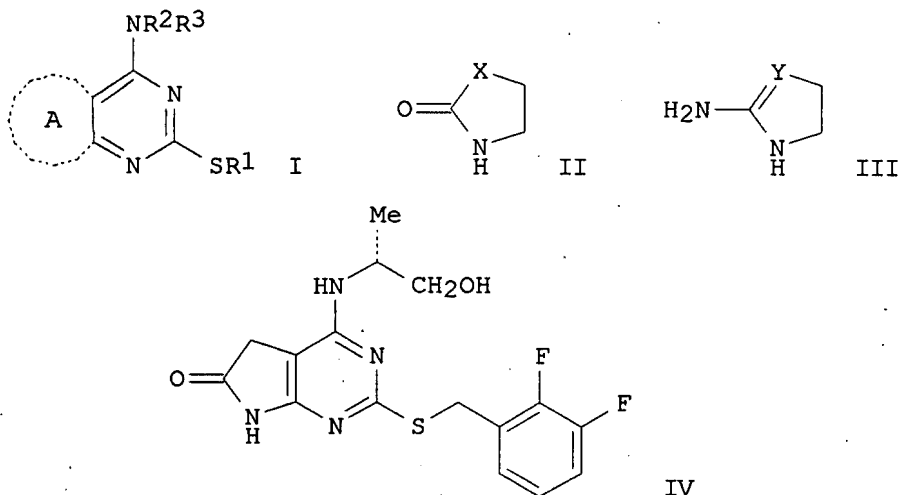
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L6 ANSWER 1 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN
ACCESSION NUMBER: 2001:597989 CAPLUS
DOCUMENT NUMBER: 135:166840
TITLE: Preparation of pyrimidine compounds as modulators of
chemokine receptor activity
INVENTOR(S): Bonnert, Roger; Cage, Peter; Hunt, Fraser; Walters,
Lain; Willis, Paul
PATENT ASSIGNEE(S): Astrazeneca Ab, Swed.
SOURCE: PCT Int. Appl., 52 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2001058902	A1	20010816	WO 2001-SE245	20010207 <--
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW			
RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG			
GB 2359078	A	20010815	GB 2000-3019	20000211 <--
EP 1265899	A1	20021218	EP 2001-902950	20010207
R:	AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR			
JP 2003522189	T	20030722	JP 2001-558051	20010207
US 2003040523	A1	20030227	US 2002-203584	20020809
US 6958344	B2	20051025		
US 2005234077	A1	20051020	US 2005-36682	20050114
PRIORITY APPLN. INFO.:			GB 2000-3019	A 20000211

OTHER SOURCE(S): MARPAT 135:166840
GI



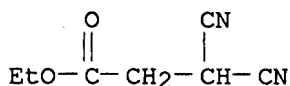
AB The title compds. [I; A = II, III (X = NH, CR₁₈R₁₉; Y = N, CR₁₈; R₁₈, R₁₉ = H, alkyl, Ph); R₁ = (un)substituted cycloalkyl, alkyl, alkenyl, etc.; R₂, R₃ = H, cycloalkyl, alkyl, etc.; NR₂R₃ = (un)substituted 3-8 membered ring optionally containing one or more atoms selected from O, S, NH, etc.], useful in treating an inflammatory disease such as psoriasis and COPD, were prepared E.g., a multi-step synthesis of the 6H-pyrrolo[2,3-d]pyrimidin-6-one IV was given. The compds. I were found to have IC₅₀ of < 10 μ M against CXCR2 receptor binding.

IT 224637-77-8

RL: RCT (Reactant); RACT (Reactant or reagent)
(preparation of pyrimidine compds. as modulators of chemokine receptor activity)

RN 224637-77-8 CAPLUS

CN Propanoic acid, 3,3-dicyano-, ethyl ester (9CI) (CA INDEX NAME)



REFERENCE COUNT: 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L6 ANSWER 2 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 2001:347933 CAPLUS

DOCUMENT NUMBER: 135:122094

TITLE: Cyclopropanation of benzylidenemalononitrile with dialkoxycarbenes and free radical rearrangement of the cyclopropanes

AUTHOR(S): Merkley, Nadine; Vennieri, Paul C.; Warkentin, John

CORPORATE SOURCE: Department of Chemistry, McMaster University, Hamilton, ON, L8S 4M1, Can.

SOURCE: Canadian Journal of Chemistry (2001), 79(3),

312-318

CODEN: CJCHAG; ISSN: 0008-4042

PUBLISHER: National Research Council of Canada

DOCUMENT TYPE: Journal

LANGUAGE: English

OTHER SOURCE(S): CASREACT 135:122094

AB Thermolysis of 2-cinnamyloxy-2-methoxy-5,5-dimethyl-Δ³-1,3,4-oxadiazoline (1a) and the analogous 2-benzyloxy-2-methoxy compound (1b) at 110°C, in benzene containing benzylidenemalononitrile, afforded products of apparent regiospecific addition of methoxycarbonyl and cinnamyl (or benzyl) radicals to the double bond. When the thermolysis of 1a was run with added TEMPO, methoxycarbonyl and cinnamyl radicals were captured. Thermolysis of the 2,2-dibenzyloxy analog (1c) in the presence of benzylidenemalononitrile gave an adduct that is formally the product of addition of benzyloxycarbonyl and benzyl radicals to the double bond. In this case, a radical addition mechanism could be ruled out, because the rate constant for decarboxylation of benzyloxycarbonyl radicals is very large. A mechanism that fits all of the results is predominant cyclopropanation of benzylidenemalononitrile by the dialkoxycarbenes derived from the oxadiazolines, in competition with fragmentation of the carbenes to radical pairs. The cyclopropanes so formed then undergo homolytic ring-opening to the appropriate diradicals. Subsequent β-scission of the diradicals to afford radical pairs, and coupling of those pairs, gives the final products. Thus, both carbene and radical chemical are involved in the overall processes.

IT 351207-62-0P 351207-63-1P 351207-65-3P

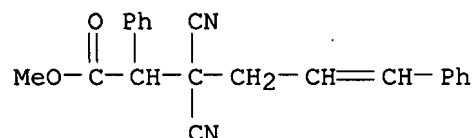
351207-66-4P 351207-67-5P

RL: SPN (Synthetic preparation); PREP (Preparation)

(cyclopropanation of benzylidenemalononitrile with dialkoxycarbenes and free radical rearrangement of the cyclopropanes)

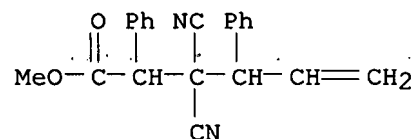
RN 351207-62-0 CAPLUS

CN Benzeneacetic acid, α-(1,1-dicyano-4-phenyl-3-butenyl)-, methyl ester (9CI) (CA INDEX NAME)



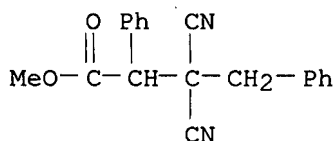
RN 351207-63-1 CAPLUS

CN Benzenebutanoic acid, β,β-dicyano-γ-ethenyl-α-phenyl-, methyl ester (9CI) (CA INDEX NAME)



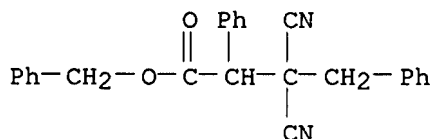
RN 351207-65-3 CAPLUS

CN Benzenebutanoic acid, β,β-dicyano-α-phenyl-, methyl ester (9CI) (CA INDEX NAME)



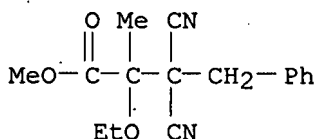
RN 351207-66-4 CAPLUS

CN Benzenebutanoic acid, β,β -dicyano- α -phenyl-, phenylmethyl ester (9CI) (CA INDEX NAME)



RN 351207-67-5 CAPLUS

CN Benzenebutanoic acid, β,β -dicyano- α -ethoxy- α -methyl-, methyl ester (9CI) (CA INDEX NAME)



REFERENCE COUNT: 20 THERE ARE 20 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L6 ANSWER 3 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 2000:310730 CAPLUS

DOCUMENT NUMBER: 133:104844

TITLE: New aspects of knoevenagel condensation and michael addition reactions on alkaline carbonates

AUTHOR(S): Aramendia, Maria A.; Borau, Victoriano; Jimenez, Cesar; Marinas, Jose M.; Romero, Francisco J.

CORPORATE SOURCE: Department of Organic Chemistry, Faculty of Sciences, Cordoba University, Cordoba, E-14004, Spain

SOURCE: Chemistry Letters (2000), (5), 574-575

CODEN: CMLTAG; ISSN: 0366-7022

PUBLISHER: Chemical Society of Japan

DOCUMENT TYPE: Journal

LANGUAGE: English

OTHER SOURCE(S): CASREACT 133:104844

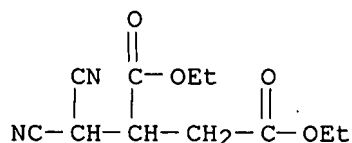
AB The Knoevenagel condensation of malononitrile with benzaldehyde on K_2CO_3 , Rb_2CO_3 and Cs_2CO_3 gave the condensation product benzylidenemalononitrile but the reaction proceeded to the hydrogenated product benzylmalononitrile. Also, the Michael addition of malononitrile to certain double bonds occurs in the presence of K_2CO_3 .

IT 82584-86-9P

RL: SPN (Synthetic preparation); PREP (Preparation)
(potassium carbonate catalyzed Michael addition reactions of malononitrile with alkenes)

RN 82584-86-9 CAPLUS

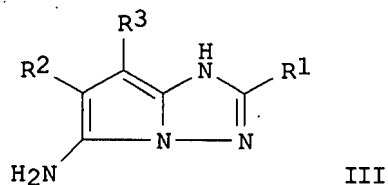
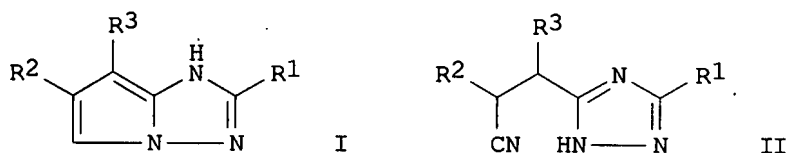
CN Butanedioic acid, (dicyanomethyl)-, diethyl ester (9CI) (CA INDEX NAME)



REFERENCE COUNT: 12 THERE ARE 12 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L6 ANSWER 4 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN
 ACCESSION NUMBER: 2000:199325 CAPLUS
 DOCUMENT NUMBER: 132:237096
 TITLE: Preparation of 1H-pyrrolo-[1,2-b][1,2,4]triazole
 INVENTOR(S): Morita, Kensuke
 PATENT ASSIGNEE(S): Fuji Photo Film Co., Ltd., Japan
 SOURCE: Jpn. Kokai Tokkyo Koho, 15 pp.
 CODEN: JKXXAF
 DOCUMENT TYPE: Patent
 LANGUAGE: Japanese
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

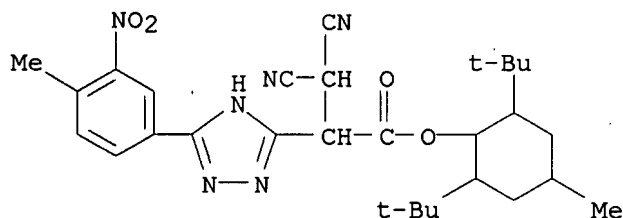
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 2000086662	A	20000328	JP 1998-265064	19980918 <--
PRIORITY APPLN. INFO.:			JP 1998-265064	19980918
OTHER SOURCE(S):			CASREACT 132:237096; MARPAT 132:237096	
GI				



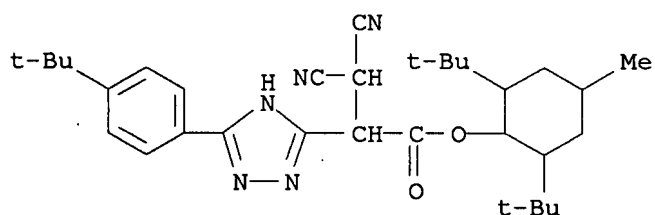
AB Title compds. I (R1-R3 = H, substituent) are prepared from triazoles II (R1-R3 = H, substituent) via III (R1-R3 = H, substituent): 3-(Tert-butylphenyl)-5-[[4-methyl-2,6-di-tert-butylhexyloxycarbonyl]bromomethyl]-1H-1,2,4-triazole was reacted with malononitrile in dimethylacetamide in the presence of NaOMe/MeOH under reflux for 30 min and reacted in the presence of CuCl in PhMe-hexane mixture under reflux for 3 h to give 91% III (R1 = 4-tert-butylphenyl; R2 = cyano, R3 = 4-methyl-2,6-di-tert-butylhexyloxycarbonyl), which was reacted with isoamyl nitrite in iso-Pr alc. at 50° for 10 h to give 40% I (R1-R3 = same as above).

IT 259266-71-2
 RL: RCT (Reactant); RACT (Reactant or reagent)
 (preparation of pyrrolotriazoles from cyanoethyltriazoles)
 RN 259266-71-2 CAPLUS

CN 1H-1,2,4-Triazole-3-acetic acid, α -(dicyanomethyl)-5-(4-methyl-3-nitrophenyl)-, 2,6-bis(1,1-dimethylethyl)-4-methylcyclohexyl ester (9CI)
(CA INDEX NAME)

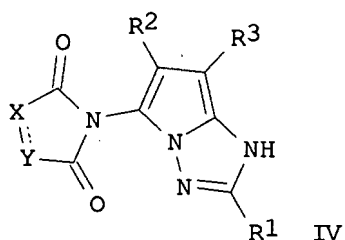
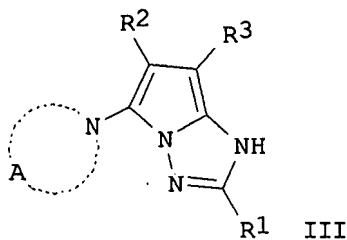
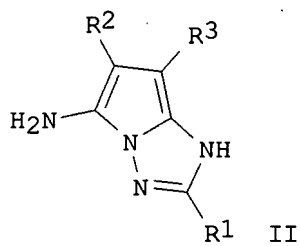
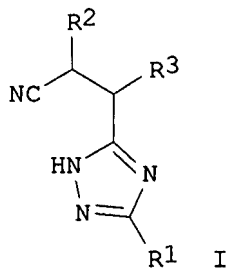


IT 259266-70-1P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation of pyrrolotriazoles from cyanoethyltriazoles)
RN 259266-70-1 CAPLUS
CN 1H-1,2,4-Triazole-3-acetic acid, α -(dicyanomethyl)-5-[4-(1,1-dimethylethyl)phenyl]-, 2,6-bis(1,1-dimethylethyl)-4-methylcyclohexyl ester (9CI) (CA INDEX NAME)

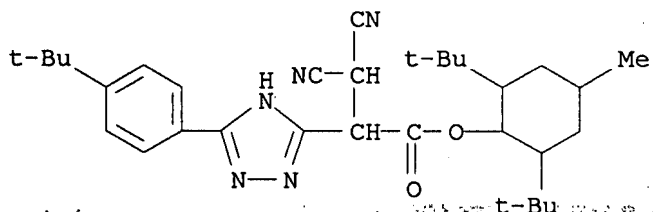


L6 ANSWER 5 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN
ACCESSION NUMBER: 2000:198039 CAPLUS
DOCUMENT NUMBER: 132:238369
TITLE: 1H-Pyrrolo[1,2-b][1,2,4]triazole derivatives and their manufacture
INVENTOR(S): Morita, Kensuke
PATENT ASSIGNEE(S): Fuji Photo Film Co., Ltd., Japan
SOURCE: Jpn. Kokai Tokkyo Koho, 14 pp.
CODEN: JKXXAF
DOCUMENT TYPE: Patent
LANGUAGE: Japanese
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 2000086661	A	20000328	JP 1998-265059	19980918 <--
PRIORITY APPLN. INFO.:			JP 1998-265059	19980918
OTHER SOURCE(S):	MARPAT 132:238369			
GI				



- AB The derivs. III and IV, useful for photog. couplers, physiol. active substances, etc., are manufactured from triazole derivs. I via intermediates II (R1-3 = H, substituent; A = non-metal atomic group to form azole ring with N; X, Y = non-metal atom to form 5-membered ring with CONCO). Thus, I (R1 = p-tert-BuC6H4; R2 = CN; R3 = 2,6-di-tert-butyl-4-methylcyclohexyloxycarbonyl) was treated with CuCl to give 91% II, 9.42 mmol of which was treated with 11.3 mmol acetylacetone in benzene in the presence of AcOH to give 8.20 mmol III (azole ring = 3,4-dimethyl-1-pyrrolyl).
- IT 259266-70-1P
 RL: IMF (Industrial manufacture); RCT (Reactant); PREP (Preparation); RACT (Reactant or reagent)
 (manufacture of pyrrolotriazole derivs.)
- RN 259266-70-1 CAPLUS
- CN 1H-1,2,4-Triazole-3-acetic acid, α -(dicyanomethyl)-5-[4-(1,1-dimethylethyl)phenyl]-, 2,6-bis(1,1-dimethylethyl)-4-methylcyclohexyl ester (9CI) (CA INDEX NAME)



L6 ANSWER 6 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN
 ACCESSION NUMBER: 2000:141061 CAPLUS
 DOCUMENT NUMBER: 132:278722
 TITLE: Spontaneous addition of active methine compounds to enol ethers and α,β -unsaturated ketones in aprotic polar solvent
 AUTHOR(S): Yokozawa, Tsutomu; Oishi, Motoi; Tanaka, Yasukazu
 CORPORATE SOURCE: Department of Applied Chemistry, Kanagawa University, Kanagawa-ku Yokohama, 221-8686, Japan
 SOURCE: Journal of Organic Chemistry (2000), 65(6),

1895-1897

CODEN: JOCEAH; ISSN: 0022-3263

PUBLISHER: American Chemical Society

DOCUMENT TYPE: Journal

LANGUAGE: English

OTHER SOURCE(S): CASREACT 132:278722

AB Addition of (EtO)2CHCH2CXYCH(CN)2 (I, X = Y = cyano; X = cyano, Y = CO2Me; X = Y = CO2Me) to enol ethers and α,β -unsatd. ketones in DMF at room temp is reported. Thus, reacting I (X = Y = cyano) with H2C:CHOEt gave (EtO)2CHCH2C(CN)2CH(OEt)Me in 63% yield. This reaction illustrates that the electron-withdrawing groups at the β -positions of the active methine group having the ones at the α and β positions were strongly affected on the acidity of I.

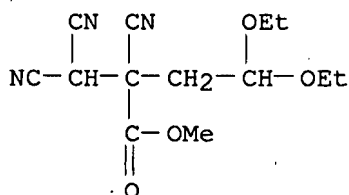
IT 184092-93-1 189348-52-5

RL: RCT (Reactant); RACT (Reactant or reagent)

(addition of methine compds. to enol ethers and α,β -unsatd. ketones)

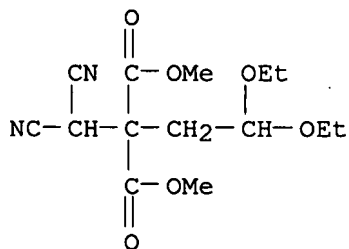
RN 184092-93-1 CAPLUS

CN Butanoic acid, 2-cyano-2-(dicyanomethyl)-4,4-diethoxy-, methyl ester (9CI)
(CA INDEX NAME)



RN 189348-52-5 CAPLUS

CN Propanedioic acid, (dicyanomethyl)(2,2-diethoxyethyl)-, dimethyl ester (9CI) (CA INDEX NAME)



IT 264142-31-6P 264142-33-8P 264142-35-0P

264142-37-2P 264142-39-4P 264142-40-7P

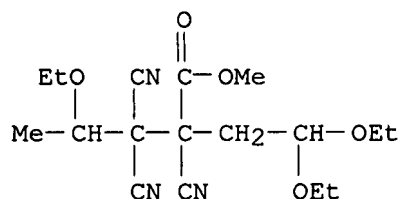
264142-41-8P 264142-43-0P 264142-45-2P

RL: SPN (Synthetic preparation); PREP (Preparation)

(addition of methine compds. to enol ethers and α,β -unsatd. ketones)

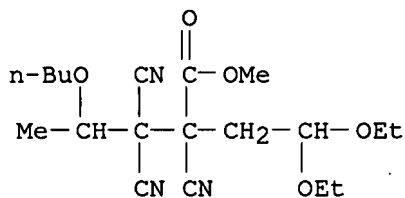
RN 264142-31-6 CAPLUS

CN Pentanoic acid, 2,3,3-tricyano-2-(2,2-diethoxyethyl)-4-ethoxy-, methyl ester (9CI) (CA INDEX NAME)



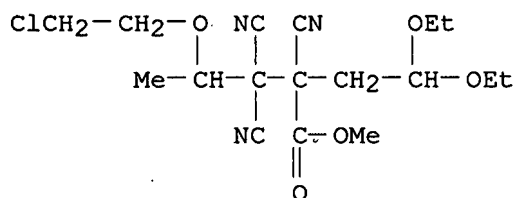
RN 264142-33-8 CAPLUS

CN Pentanoic acid, 4-butoxy-2,3,3-tricyano-2-(2,2-diethoxyethyl)-, methyl ester (9CI) (CA INDEX NAME)



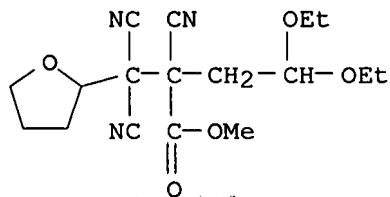
RN 264142-35-0 CAPLUS

CN Pentanoic acid, 4-(2-chloroethoxy)-2,3,3-tricyano-2-(2,2-diethoxyethyl)-, methyl ester (9CI) (CA INDEX NAME)



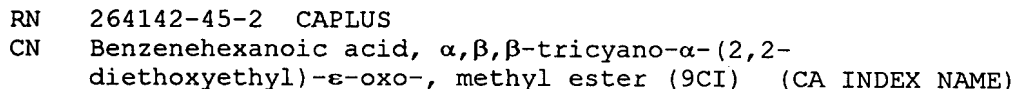
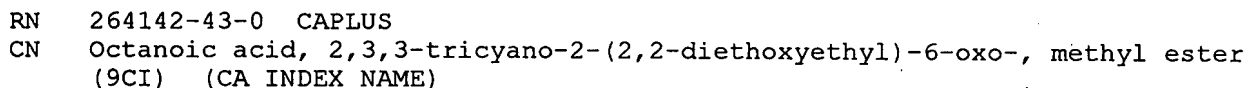
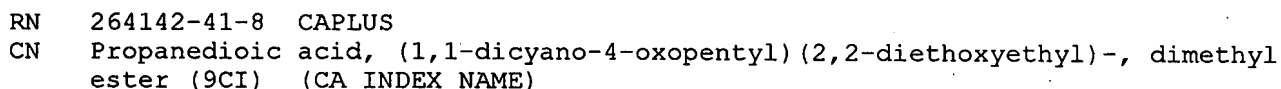
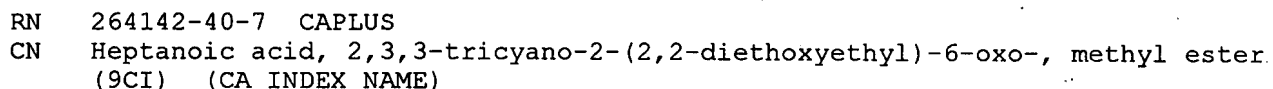
RN 264142-37-2 CAPLUS

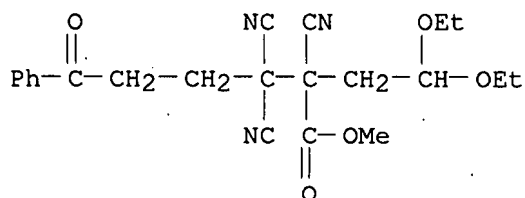
CN 2-Furanpropanoic acid, α,β,β -tricyano- α -(2,2-diethoxyethyl)tetrahydro-, methyl ester (9CI) (CA INDEX NAME)



RN 264142-39-4 CAPLUS

CN 2H-Pyran-2-propanoic acid, α,β,β -tricyano- α -(2,2-diethoxyethyl)tetrahydro-, methyl ester (9CI) (CA INDEX NAME)

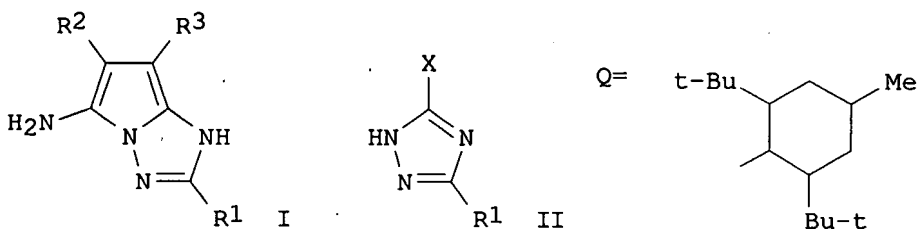




REFERENCE COUNT: 25 THERE ARE 25 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L6 ANSWER 7 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN
 ACCESSION NUMBER: 2000:139178 CAPLUS
 DOCUMENT NUMBER: 132:180579
 TITLE: Preparation of 1H-pyrrolo[1,2-b][1,2,4]triazol-5-ylamines
 INVENTOR(S): Morita, Kensuke
 PATENT ASSIGNEE(S): Fuji Photo Film Co., Ltd., Japan
 SOURCE: Jpn. Kokai Tokkyo Koho, 15 pp.
 CODEN: JKXXAF
 DOCUMENT TYPE: Patent
 LANGUAGE: Japanese
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 2000063382	A	20000229	JP 1998-232925	19980819 <--
PRIORITY APPLN. INFO.:			JP 1998-232925	19980819
OTHER SOURCE(S):		CASREACT 132:180579; MARPAT 132:180579		
GI				

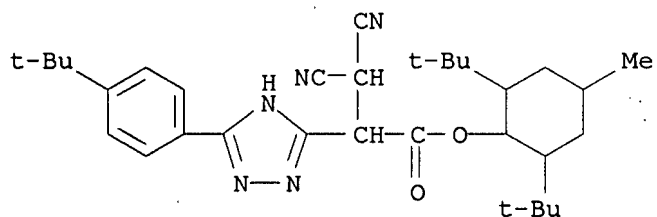


AB Title compds. I (R¹-R³ = H, substituent), useful as intermediates for physiol. active substances, photog. couplers, dyes, etc., are prepared from triazoles II (X = CHR³CHR²CN; R¹-R³ = same as I). II (R¹ = C₆H₄Bu-t-p, X = CHBrCO₂Q) was treated with malononitrile and MeONa in DMF-MeOH under ice-cooling for 30 min and heated in the presence of CuCl in PhMe-hexane under reflux for 3 h to give 91% I (R¹ = C₆H₄Bu-t-p, R² = cyano, R³ = CO₂Q).

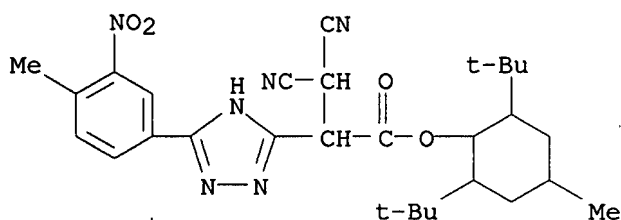
IT 259266-70-1P 259266-71-2P
 RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
 (preparation of pyrrolotriazolylamines as intermediates for physiol. active substances, dyes, and photog. couplers)

RN 259266-70-1 CAPLUS

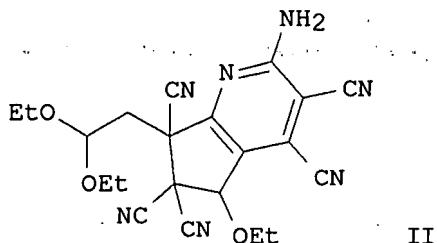
CN 1H-1,2,4-Triazole-3-acetic acid, α-(dicyanomethyl)-5-[4-(1,1-dimethylethyl)phenyl]-, 2,6-bis(1,1-dimethylethyl)-4-methylcyclohexyl ester (9CI) (CA INDEX NAME)



RN 259266-71-2 CAPLUS
 CN 1H-1,2,4-Triazole-3-acetic acid, α -(dicyanomethyl)-5-(4-methyl-3-nitrophenyl)-, 2,6-bis(1,1-dimethylethyl)-4-methylcyclohexyl ester (9CI)
 (CA INDEX NAME)



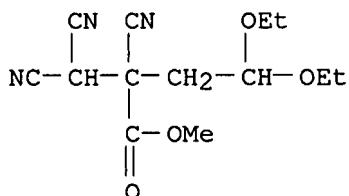
L6 ANSWER 8 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN
 ACCESSION NUMBER: 1999:380253 CAPLUS
 DOCUMENT NUMBER: 131:170252
 TITLE: Tandem dimerization and double annulation of 3,3,4,4-tetracyanobutanal acetal. Synthesis of a bicyclic 2-aminopyridine derivative
 AUTHOR(S): Yokozawa, Tsutomu; Nishikata, Akira; Kimura, Takamasa; Shimizu, Kazuki; Takehana, Tomoyuki
 CORPORATE SOURCE: Department of Applied Chemistry, Kanagawa University, Yokohama, 221-8686, Japan
 SOURCE: Tetrahedron Letters (1999), 40(25), 4707-4710
 CODEN: TELEAY; ISSN: 0040-4039
 PUBLISHER: Elsevier Science Ltd.
 DOCUMENT TYPE: Journal
 LANGUAGE: English
 OTHER SOURCE(S): CASREACT 131:170252
 GI



AB 3,3,4,4-Tetracyanobutanal acetal (I), which is easily obtained from tetracyanoethylene, Et vinyl ether, and ethanol, yielded 2-aminopyridine derivative II fused with cyclopentane in one pot in the presence of pyridine.

On the basis of several expts., the proposed mechanism involves the Michael reaction of I with the diene generated by the elimination of hydrogen cyanide and ethanol from I, followed by double intramol. nucleophilic addns. to the cyano groups.

IT 184092-93-1
 RL: RCT (Reactant); RACT (Reactant or reagent)
 (preparation of bicyclic aminopyridine by tandem dimerization-cyclization of tetracyanobutanol acetal)
 RN 184092-93-1 CAPLUS
 CN Butanoic acid, 2-cyano-2-(dicyanomethyl)-4,4-diethoxy-, methyl ester (9CI)
 (CA INDEX NAME)



REFERENCE COUNT: 13 THERE ARE 13 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L6 ANSWER 9 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1999:142792 CAPLUS

DOCUMENT NUMBER: 130:360507

TITLE: An N2S2 Bifunctional Chelator for Technetium-99m and Rhenium: Complexation, Conjugation, and Epimerization to a Single Isomer

AUTHOR(S): Luyt, Leonard G.; Jenkins, Hilary A.; Hunter, Duncan H.

CORPORATE SOURCE: Department of Chemistry, University of Western Ontario, London, ON, N6A 5B7, Can.

SOURCE: Bioconjugate Chemistry (1999), 10(3), 470-479

CODEN: BCCHES; ISSN: 1043-1802

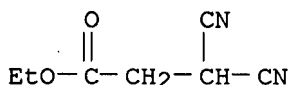
PUBLISHER: American Chemical Society

DOCUMENT TYPE: Journal

LANGUAGE: English

AB A bifunctional chelator HO₂CCH(CH₂NHCOCH₂SH)₂ 6 (H₃L) was prepared bearing an N₂S₂ core for binding Re or Tc and a carboxylic acid group for conjugation to amino groups of biomols. Complexation of 6 with Re(V) resulted in two kinetic isomers, anti- and syn-[ReO(HL)]- 7, being formed in approx. equal amts. Epimerization with 0.5M NaOH yields a single isomer anti-7, as determined by NMR spectroscopy and single-crystal x-ray anal. [99mTcO(HL)]- was prepared at the tracer level by reaction of the ligand with 99mTcO₄-, SnCl₂ and Na gluconate giving a mixture of two isomers, but showing a preference for the anti isomer. Chelation in the presence of 1 M NaOH results in anti-8 being formed as the sole product. The bifunctional ability of the ligand was explored by amide formation with (S)-α-phenethylamine, either by direct DCC coupling or through the RO₂CCH(CH₂NHCOCH₂S^{Tr})₂ 9 (R = succinimidyl) intermediate. The deprotected bioconjugate PhCHMeNHOCCH(CH₂NHCOCH₂SH)₂ 11 (H₂L1) was complexed with Re, yielding similar amts. of two isomeric Re complexes, anti- and syn-12, which were isolated and characterized by NMR spectroscopy. Treatment of the kinetic mixture of anti- and syn-[ReOL1]- 12 with 1 M NaOH resulted in quant. conversion to a single Re complex anti-12. With 99mTc in 0.1M NaOAc, bioconjugate 11 yielded anti- and sym-[99mTcOL1]- 13 in a 2:1 ratio, resp. In contrast, complexation in the presence of 1 M NaOH gave only one 99mTc complex, assigned the structure anti-13.

IT 224637-77-8P
 RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
 (Reactant or reagent)
 (preparation and reactant for preparation of rhenium and technetium
 bis(thioacetamidomethyl)propionate and bis(thioacetamidomethyl)propanam
 ide complexes)
 RN 224637-77-8 CAPLUS
 CN Propanoic acid, 3,3-dicyano-, ethyl ester (9CI) (CA INDEX NAME)



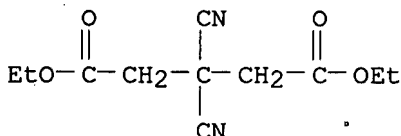
REFERENCE COUNT: 24 THERE ARE 24 CITED REFERENCES AVAILABLE FOR THIS
 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L6 ANSWER 10 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN
 ACCESSION NUMBER: 1999:142376 CAPLUS
 DOCUMENT NUMBER: 130:239567
 TITLE: Diazaspirononanium salt for use as template for
 zeolite synthesis
 INVENTOR(S): Kubota, Yoshihiro; Sugi, Yoshihiro
 PATENT ASSIGNEE(S): Showa Denko K. K., Japan
 SOURCE: Jpn. Kokai Tokkyo Koho, 9 pp.
 CODEN: JKXXAF
 DOCUMENT TYPE: Patent
 LANGUAGE: Japanese
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 11060577	A	19990302	JP 1997-220414	19970815 <--
PRIORITY APPLN. INFO.:			JP 1997-220414	19970815
OTHER SOURCE(S):	MARPAT 130:239567			

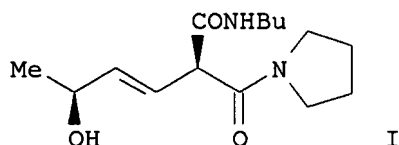
AB Claimed template is a salt of substituted 2,7-diazaspiro[4,4]nonanium.
 Hydrothermal synthesis of a zeolite by bringing a silica source and/or an
 alumina source into contact with the zeolite is also claimed. ZSM-12
 zeolites having crystal size of a major axis $\geq 50 \mu\text{m}$ are also
 claimed.

IT 77415-69-1P
 RL: IMF (Industrial manufacture); RCT (Reactant); PREP (Preparation); RACT
 (Reactant or reagent)
 (reaction of; diazaspirononanium salts as templates for manufacture of
 ZSM-12 zeolites having large crystal size)
 RN 77415-69-1 CAPLUS
 CN Pentanedioic acid, 3,3-dicyano-, diethyl ester (9CI) (CA INDEX NAME)



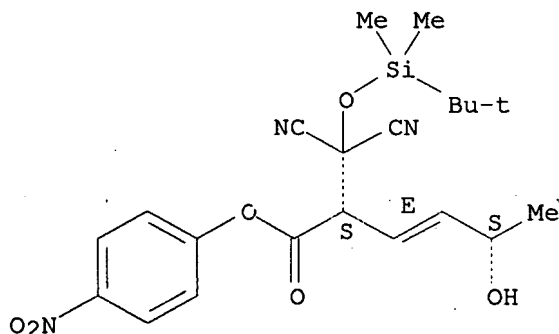
L6 ANSWER 11 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN
 ACCESSION NUMBER: 1999:118566 CAPLUS
 DOCUMENT NUMBER: 130:237421

TITLE: Stereoselective synthesis of alkenylated malonic diamide
 AUTHOR(S): Nemoto, Hisao; Ibaragi, Touru; Bando, Masahiko; Kido, Masaru; Shibuya, Masayuki
 CORPORATE SOURCE: Faculty of Pharmaceutical Sciences, the University of Tokushima, Tokushima, 770-8505, Japan
 SOURCE: Tetrahedron Letters (1999), 40(7), 1319-1322
 CODEN: TELEAY; ISSN: 0040-4039
 PUBLISHER: Elsevier Science Ltd.
 DOCUMENT TYPE: Journal
 LANGUAGE: English
 OTHER SOURCE(S): CASREACT 130:237421
 GI



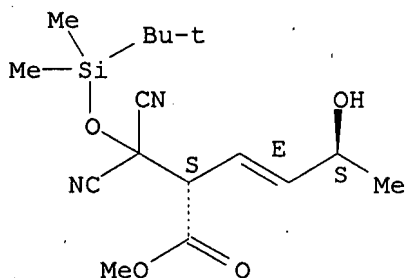
AB A highly stereoselective synthesis of an alkenylated malonic diamide (I) starting from a γ,δ -epoxy α,β -unsatd. carboxamide was accomplished using masked acyl cyanide (protected hydroxymalonitrile) via palladium-catalyzed regio- and stereoselective carbon-carbon bond formation.
 IT 221219-72-3P 221219-73-4P
 RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
 (stereoselective synthesis of alkenylated malonic diamide using masked acyl cyanide)
 RN 221219-72-3 CAPLUS
 CN 3-Hexenoic acid, 2-[dicyano[[(1,1-dimethylethyl)dimethylsilyl]oxy]methyl]-5-hydroxy-, 4-nitrophenyl ester, (2R,3E,5R)-rel- (9CI) (CA INDEX NAME)

Relative stereochemistry.
 Double bond geometry as shown.



RN 221219-73-4 CAPLUS
 CN 3-Hexenoic acid, 2-[dicyano[[(1,1-dimethylethyl)dimethylsilyl]oxy]methyl]-5-hydroxy-, methyl ester, (2R,3E,5R)-rel- (9CI) (CA INDEX NAME)

Relative stereochemistry.
 Double bond geometry as shown.



REFERENCE COUNT: 13 THERE ARE 13 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L6 ANSWER 12 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1998:402031 CAPLUS

DOCUMENT NUMBER: 129:122635

TITLE: The reaction of phenacylmalononitrile with hydrazines: synthesis of new pyridazinones and pyrazolo[1,5-a]pyrimidines

AUTHOR(S): Elnagdi, Mohamed Hilmy; El-Ghamry, Ibrahim; Kandeel, Ezz; Abdel Rahman, A. H.; Al-Naggar, Abdul Aziz; Amer, Samir; Riad, Mohamed

CORPORATE SOURCE: Department of Chemistry, Faculty of Science, University of Kuwait, Safat, 13060, Kuwait

SOURCE: Gazzetta Chimica Italiana (1997), 127(12), 791-794

CODEN: GCITA9; ISSN: 0016-5603

PUBLISHER: Societa Chimica Italiana

DOCUMENT TYPE: Journal

LANGUAGE: English

OTHER SOURCE(S): CASREACT 129:122635

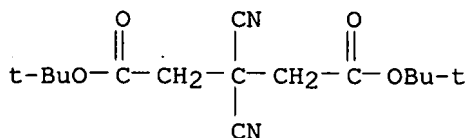
AB The reaction of phenacylmalononitrile with hydrazine hydrate affords a mixture of 3,5-diamino-4-phenacylpyrazole (I), and 6-oxo-3-phenyl-1,4,5,6-tetrahydropyridazine-5-carbonitrile. The reaction of I with a variety of reagents, that enabled the synthesis of some new pyrazolo[1,5-a]pyrimidine derivs., is described.

IT 210347-41-4P

RL: SPN (Synthetic preparation); PREP (Preparation)
(synthesis of pyridazinones and pyrazolo[1,5-a]pyrimidines)

RN 210347-41-4 CAPLUS

CN Pentanedioic acid, 3,3-dicyano-, bis(1,1-dimethylethyl) ester (9CI) (CA INDEX NAME)



REFERENCE COUNT: 15 THERE ARE 15 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L6 ANSWER 13 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1998:76233 CAPLUS

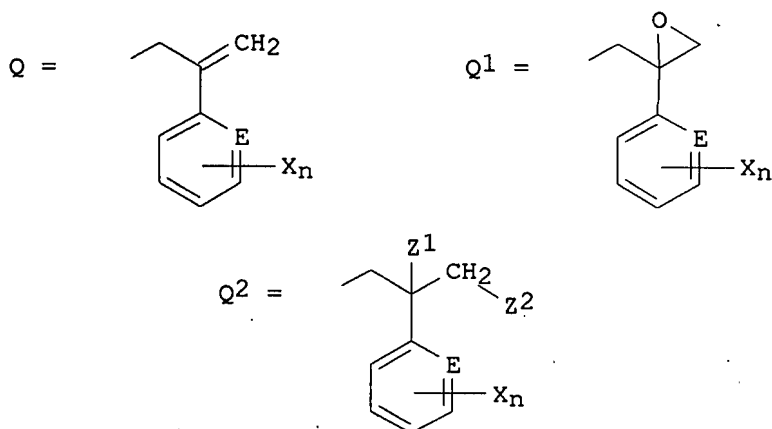
DOCUMENT NUMBER: 128:177233

TITLE: Malononitrile derivatives and herbicides containing them

INVENTOR(S): Hosokawa, Akemi; Ikeda, Osamu
 PATENT ASSIGNEE(S): Mitsubishi Chemical Industries Ltd., Japan
 SOURCE: Jpn. Kokai Tokkyo Koho, 18 pp.
 CODEN: JKXXAF
 DOCUMENT TYPE: Patent
 LANGUAGE: Japanese
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 10029966	A	19980203	JP 1996-187796	19960717 <--
PRIORITY APPLN. INFO.:			JP 1996-187796	19960717
OTHER SOURCE(S):	MARPAT	128:177233		

GI



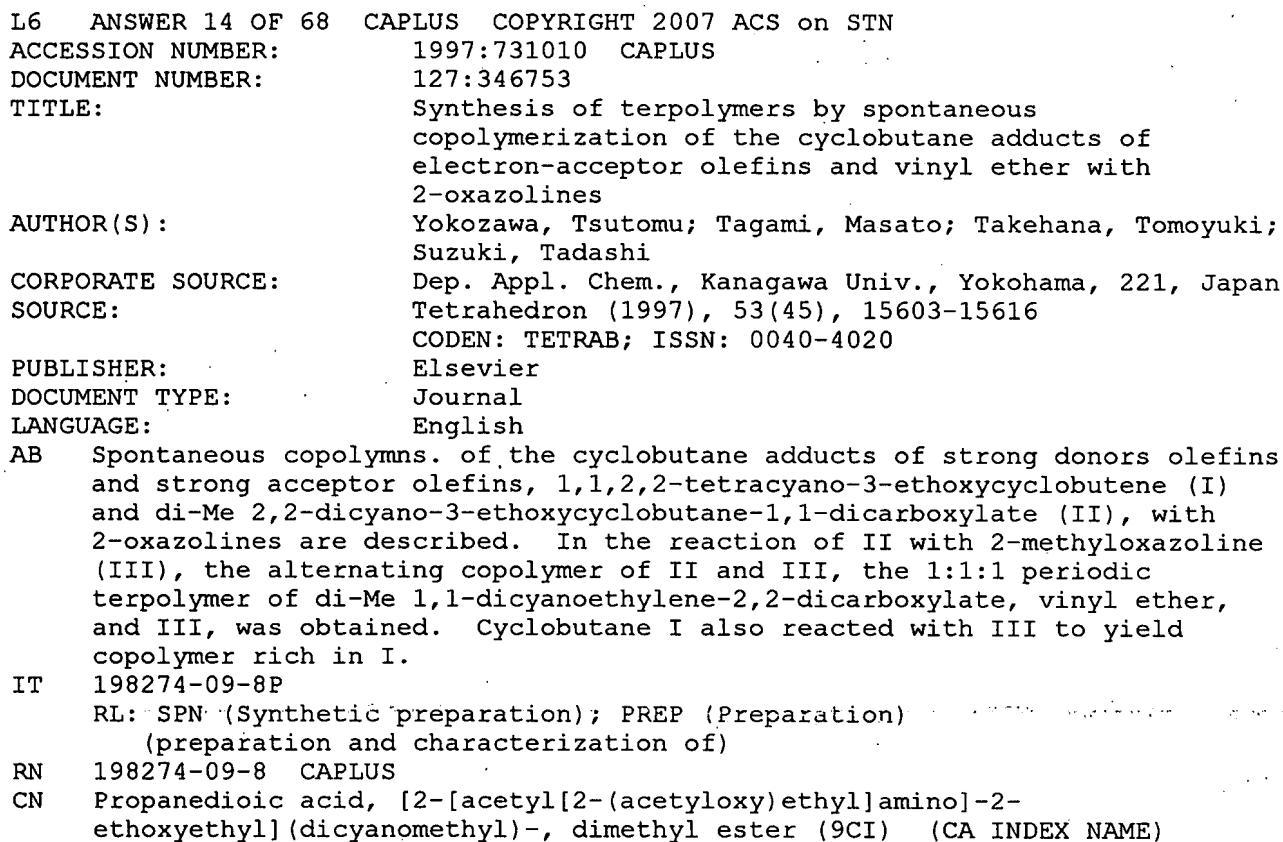
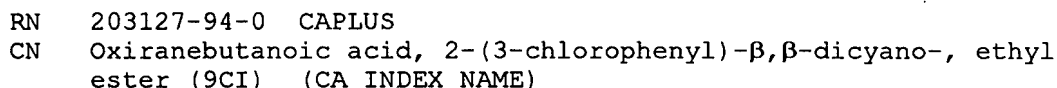
AB The derivs. are represented by R1R2C(CN)2 [I; R1 = H, C1-6 alkyl, C4-7 cycloalkyl, C2-6 alkenyl, C2-6 alkynyl, C1-6 haloalkyl, C2-6 haloalkenyl, C2-6 haloalkynyl, C2-8 alkoxyalkyl, C3-6 alkoxyalkyl, C2-4 cyanoalkyl, C1-6 hydroxyalkyl, C2-7 alkylamido, C7-9 aralkyl, C8-12 arylcarbonylalkyl, (un)substituted Ph, (un)substituted pyridyl, (un)substituted thiazolyl, CR3R4A; A = (un)substituted Ph, pyridyl, thiazolyl; R3-4 = H, Me; R2 = Q, Q1, Q2; E = CH, N; X = halo, C1-4 alkyl, C1-3 haloalkyl, NO2, C1-8 haloalkoxy, (un)substituted benzyloxy, pyridyloxy; n = 0-2; Z1-2 = OH, halo, C1-4 alkylsulfonyloxy, (un)substituted phenylsulfonyloxy]. The herbicides contain I as active ingredients. I (R1 = H, R2 = Q, E = CH, X = 3-Me, 5-Me) showed 91-100% herbicidal activity against Echinochloa oryzicola, Monochoria vaginalis, and Scirpus juncoides.

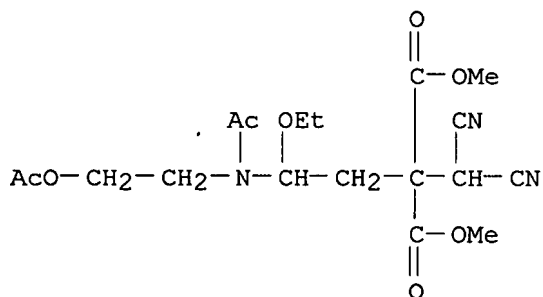
IT 203127-60-0P 203127-94-0P

RL: AGR (Agricultural use); BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation); USES (Uses)
 (preparation of malononitrile derivs. as herbicides)

RN 203127-60-0 CAPLUS

CN Benzenepentanoic acid, 3-chloro-β,β-dicyano-δ-methylene-, ethyl ester (9CI) (CA INDEX NAME)





REFERENCE COUNT: 37 THERE ARE 37 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L6 ANSWER 15 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1997:513581 CAPLUS

DOCUMENT NUMBER: 127:184884

TITLE: Multinuclear cluster complexes as diagnostic imaging contrast agents

INVENTOR(S): Droege, Michael; Yu, Shi-Bao; Sanderson, William; Bacon, Edward; Delecki, Daniel; Earley, William; Ye, Naidong

PATENT ASSIGNEE(S): Nycomed Salutar, Inc., USA

SOURCE: PCT Int. Appl., 51 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 9726921	A2	19970731	WO 1997-GB211	19970123 <--
WO 9726921	A3	19971023		
W: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, HU, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, TJ, TM, TR, TT, UA, UG, US, UZ, VN				
RW: KE, LS, MW, SD, SZ, UG, AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG				
CA 2241190	A1	19970731	CA 1997-2241190	19970123 <--
AU 9714523	A	19970820	AU 1997-14523	19970123 <--
EP 876161	A2	19981111	EP 1997-901180	19970123 <--
EP 876161	B1	20011121		
R: DE, DK, ES, FR, GB, IT, SE, IE, FI				
CN 1208353	A	19990217	CN 1997-191757	19970123 <--
HU 9901488	A2	19990830	HU 1999-1488	19970123 <--
BR 9707300	A	19991228	BR 1997-7300	19970123 <--
JP 2000515850	T	20001128	JP 1997-526668	19970123 <--
NO 9803371	A	19980722	NO 1998-3371	19980722 <--
PRIORITY APPLN. INFO.:			GB 1996-1340	A 19960123
			WO 1997-GB211	W 19970123

OTHER SOURCE(S): MARPAT 127:184884

AB Diagnostic imaging contrast media are claimed comprising a physiologically tolerable image contrast-enhancing complex, said complex comprising a pair of interconjugated multinuclear clusters, together with at least one pharmaceutical carrier or excipient. Included, for example, are multinuclear cluster complexes (M₃)₂L₃ containing three metal atoms and L is a ligand. Clusters (M₃)₂L₃ include M₃ = M₃SO_b where a = 1-4, b = 0-3 and a + b = 4, e.g., M₃ = W₃SO₃. Ligands L include various

polyaminocarboxylates and derivs. represented by general formula (R2)2N[(CHR4)mNR1]n(CHR4)mN(R2)2, e.g., N'-serinol-, N'-methyl-, and N'-benzyl-N,N,N',N''-diethylenetriaminetetraacetic acids, various N'-(polyhydroxyalkyl)-N'-methyldiethylenetriaminetetraacetic acids, 2-carboxymethylpropylenediaminetetraacetic acid, etc., for which preps. are given of these and other example ligands. Preparation of cluster compds., e.g., Na4[(W3SO3)2(EGTA)3] (EGTA = ethyleneglycol bis(2-aminoethyl ether)-N,N,N',N''-tetraacetate), from [W3SO3(H2O)9]Cl4 and the appropriate polyaminocarboxylic acid ligand, are described. The claimed preparation of [W3SO3(H2O)9]Cl4 comprises reaction of W(CO)6 and Na2S, followed by acidification of the product with at least 6 N HCl, and purification. A charged contrast medium complex may be post-complexed with, e.g., choline hydrochloride or N-methyl-N,N-bis(hydroxyethyl)ethylenediamine, to give a preferred neutral derivative. Pharmaceutically acceptable forms of the diagnostic imaging contrast media comprising said cluster complexes and dosages for the x-ray contrast media are briefly discussed.

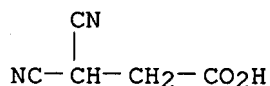
IT 194083-97-1P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(for preparation of multinuclear tungsten aminocarboxylate cluster complexes as diagnostic imaging contrast agents)

RN 194083-97-1 CAPLUS

CN Propanoic acid, 3,3-dicyano- (9CI) (CA INDEX NAME)



L6 ANSWER 16 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1997:414890 CAPLUS

DOCUMENT NUMBER: 127:144690

TITLE: Metabolism and disposition of the antifolate LY231514 in mice and dogs

AUTHOR(S): Woodland, J. M.; Barnett, C. J.; Dorman, D. E.; Gruber, J. M.; Shih, C.; Spangle, L. A.; Wilson, T. M.; Ehlhardt, W. J.

CORPORATE SOURCE: Lilly Res. Laboratories, USA

SOURCE: Drug Metabolism and Disposition (1997), 25(6), 693-700

CODEN: DMDSAI; ISSN: 0090-9556

PUBLISHER: Williams & Wilkins

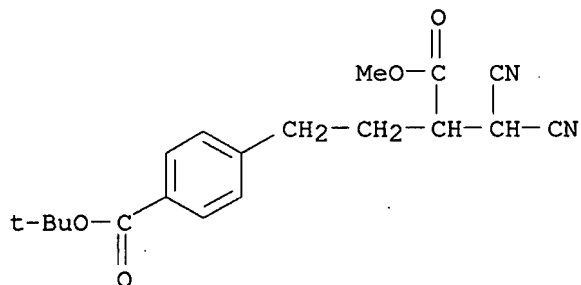
DOCUMENT TYPE: Journal

LANGUAGE: English

AB The metabolism and disposition of LY231514 was studied in mice and dogs. LY231514 is a novel pyrrolopyrimidine-based multi-target antifolate (MTA) showing broad in vivo antitumor activity in mouse models and is currently in phase II human clinical trials. Doses (i.v.) of the compound showed high plasma levels, resulting in AUC values of 30-33 µg-hr/mL for mice and dogs after 20 and 7.5 mg/kg doses, resp. The compound was eliminated rapidly. Half-life values for mice and dogs were about 7 and 2 h, resp. In vitro plasma binding measured 56% in mice, 46% in dogs, and 81% in humans. Fecal elimination was the major excretion pathway in mice after single i.v. doses of [14C]LY231514. Urine constituted the major route of excretion in dogs. Parent LY231514 accounted for the majority of urinary radiocarbon in mice (90%) and dogs (68%). Minor metabolites were found in urine, but the amounts were too small to isolate or identify. Based on an earlier observation that LY231514 photodegraded to produce reaction products having similar retention times as these minor urinary isolates, a photo oxidation system was developed which in fact produced these

metabolites. Subsequently, these photolytically produced materials were used as stds. to identify two novel in vivo metabolites formed by oxidation of the pyrrolo-pyrimidine ring system of LY231514. The oxidative transformations are similar to those observed for tryptophan and other indoles in that the pyrrole ring is oxidized to give an amide; further oxidation cleaves this ring, one ring carbon is lost, and a ketone is formed.

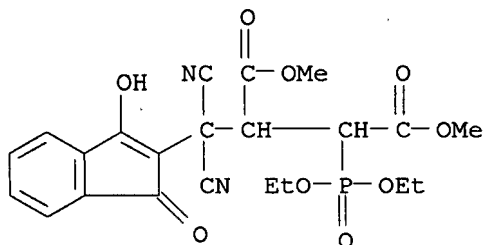
IT 193265-49-5P
 RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
 (antifolate drug LY231514 metabolism and pharmacokinetics in mice and dogs)
 RN 193265-49-5 CAPLUS
 CN Benzenebutanoic acid, α -(dicyanomethyl)-4-[(1,1-dimethylethoxy)carbonyl]-, methyl ester (9CI) (CA INDEX NAME)



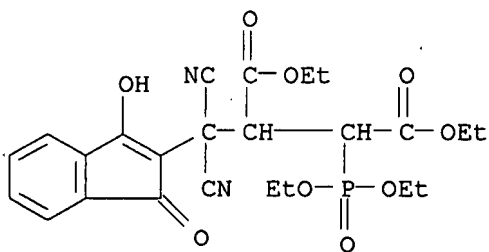
L6 ANSWER 17 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN
 ACCESSION NUMBER: 1997:330734 CAPLUS
 DOCUMENT NUMBER: 127:34293
 TITLE: The reactions of Wittig-Horner reagents with 1,3-dioxo- Δ^2 , α -indanmalononitrile
 AUTHOR(S): Boulos, Leila Sadek; Yakout, El-Sayed M. A.
 CORPORATE SOURCE: National Research Centre, Cairo, Egypt
 SOURCE: Heteroatom Chemistry (1997), 8(3), 253-257
 CODEN: HETCE8; ISSN: 1042-7163
 PUBLISHER: Wiley
 DOCUMENT TYPE: Journal
 LANGUAGE: English

AB Wittig-Horner reagents react with 1,3-dioxo- Δ^2 , α -indanmalononitrile to give phosphonate adducts. Structural reasoning for the new products was based on compatible anal. and spectral data (IR, ^1H , ^{31}P NMR, and MS). The mechanism that accounts for the formation of the new adducts is discussed.

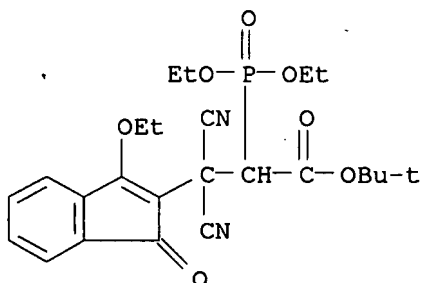
IT 190722-21-5P 190722-23-7P 190722-25-9P
 RL: SPN (Synthetic preparation); PREP (Preparation)
 (preparation of)
 RN 190722-21-5 CAPLUS
 CN Butanedioic acid, 2-[dicyano(3-hydroxy-1-oxo-1H-inden-2-yl)methyl]-3-(diethoxyphosphinyl)-, dimethyl ester (9CI) (CA INDEX NAME)



RN 190722-23-7 CAPLUS
 CN Butanedioic acid, 2-[dicyano(3-hydroxy-1-oxo-1H-inden-2-yl)methyl]-3-(diethoxyphosphinyl)-, diethyl ester (9CI) (CA INDEX NAME)



RN 190722-25-9 CAPLUS
 CN 1H-Indene-2-propanoic acid, β,β -dicyano- α -(diethoxyphosphinyl)-3-ethoxy-1-oxo-, 1,1-dimethylethyl ester (9CI) (CA INDEX NAME)



REFERENCE COUNT: 18 THERE ARE 18 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L6 ANSWER 18 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1997:274825 CAPLUS

DOCUMENT NUMBER: 126:317775

TITLE: Ring-opening polymerization of cyclobutane adduct of dimethyl 1,1-dicyanoethylene-2,2-dicarboxylate and ethyl vinyl ether

AUTHOR(S): Yokozawa, Tsutomu; Wakabayashi, Yuki; Kimura, Takamasa
 CORPORATE SOURCE: Dep. Applied Chem., Kanagawa Univ., Yokohama, 221, Japan

SOURCE: Journal of Polymer Science, Part A: Polymer Chemistry (1997), 35(8), 1563-1570
 CODEN: JPACEC; ISSN: 0887-624X

PUBLISHER: Wiley

DOCUMENT TYPE: Journal

LANGUAGE: English

AB For an extension of the work on the ring-opening polymns. of cyclobutane adducts of strong donor olefins and strong acceptor olefins yielding novel alternating copolymers of those olefins, the ring-opening polymerization of the cyclobutane adduct (I; di-Me 2,2-dicyano-3-ethoxy-1,1-cyclobutanedicarboxylate) of di-Me 1,1-dicyanoethylene-2,2-dicarboxylate (DDED) and Et vinyl ether (EVE) is investigated. I reacted with methanol and acetic acid at ambient temperature to yield the corresponding ring-opened adducts. Polymns. of I were carried out with anionic initiators, tertiary amines, ammonium halides, and Lewis acids, resp., according to the polymerization

methods of the cyclobutane adduct of tetracyanoethylene and EVE. All these polymerization catalysts except for ammonium halides were effective for the polymerization of I, yielding alternating copolymers of DDED and EVE. The chain

transfer reactions of the polymerization with anionic initiators are also discussed on the basis of a model reaction.

IT 189348-52-5

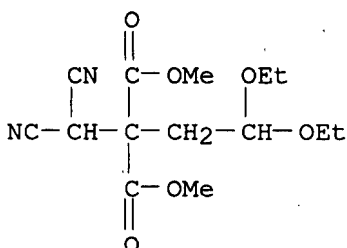
RL: RCT (Reactant); RACT (Reactant or reagent)

(model reaction for determination of mechanism; ring-opening polymerization of di-Me

2,2-dicyano-3-ethoxy-1,1-cyclobutanedicarboxylate)

RN 189348-52-5 CAPLUS

CN Propanedioic acid, (dicyanomethyl) (2,2-diethoxyethyl)-, dimethyl ester (9CI) (CA INDEX NAME)



IT 189348-50-3P 189348-51-4P

RL: SPN (Synthetic preparation); PREP (Preparation)

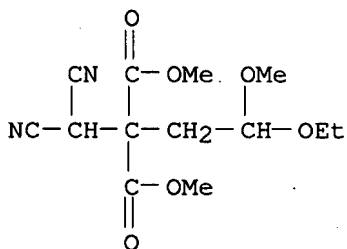
(model reactions for polymerization; reactions of di-Me

2,2-dicyano-3-ethoxy-

1,1-cyclobutanedicarboxylate with acetic acid and methanol)

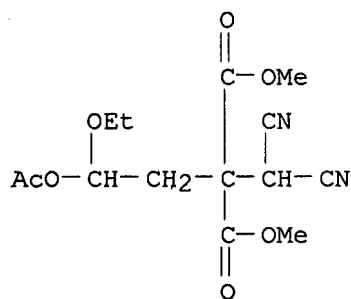
RN 189348-50-3 CAPLUS

CN Propanedioic acid, (dicyanomethyl) (2-ethoxy-2-methoxyethyl)-, dimethyl ester (9CI) (CA INDEX NAME)

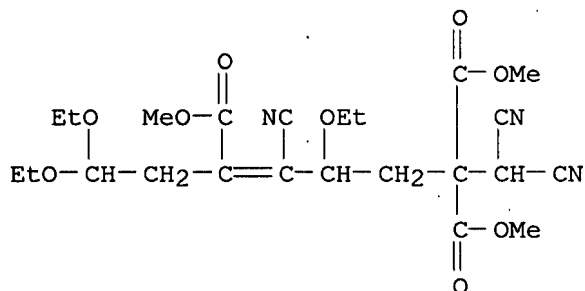


RN- 189348-51-4 CAPLUS

CN Propanedioic acid, [2-(acetyloxy)-2-ethoxyethyl] (dicyanomethyl)-, dimethyl ester (9CI) (CA INDEX NAME)



IT 189348-53-6P
 RL: SPN (Synthetic preparation); PREP (Preparation)
 (ring-opening polymerization of di-Me 2,2-dicyano-3-ethoxy-1,1-cyclobutanedicarboxylate)
 RN 189348-53-6 CAPLUS
 CN 5-Octene-2,2,6-tricarboxylic acid, 1,1,5-tricyano-4,8,8-triethoxy-, trimethyl ester (9CI) (CA INDEX NAME)



REFERENCE COUNT: 27 THERE ARE 27 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L6 ANSWER 19 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1996:672685 CAPLUS

DOCUMENT NUMBER: 126:8777

TITLE: Ring-Opening Polymerization of the Cyclobutane Adduct of Methyl Tricyanoethylenecarboxylate and Ethyl Vinyl Ether

AUTHOR(S): Yokozawa, Tsutomu; Tsuruta, Ei-ichi

CORPORATE SOURCE: Department of Applied Chemistry, Kanagawa University, Yokohama, 221, Japan

SOURCE: Macromolecules (1996), 29(25), 8053-8056

CODEN: MAMOBX; ISSN: 0024-9297

PUBLISHER: American Chemical Society

DOCUMENT TYPE: Journal

LANGUAGE: English

AB The ring-opening polymns. of a cyclobutane adduct (I) of Me tricyanoethylenecarboxylate (MTCE) and Et vinyl ether (EVE) are investigated. The adduct I reacted with acetic acid and ethanol at ambient temperature to yield the ring-opened corresponding adducts in good yields. I was polymerized with Lewis acids, anionic initiators, tertiary amines, and ammonium halides. All the catalysts except for ammonium halides were effective for the alternating polymerization similar to the polymerization of cyclobutane adduct of TCNE and EVE.

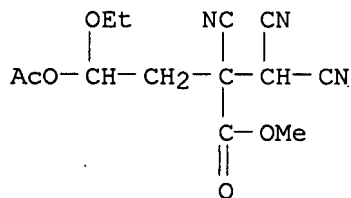
IT 184092-92-0P 184092-93-1P

RL: SPN (Synthetic preparation); PREP (Preparation)
 (ring-opening reactivity of cyclobutane adduct of Me

tricyanoethylenecarboxylate with ethanol or acetic acid and Et vinyl ether)

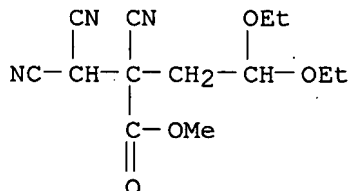
RN 184092-92-0 CAPLUS

CN Butanoic acid, 4-(acetyloxy)-2-cyano-2-(dicyanomethyl)-4-ethoxy-, methyl ester (9CI) (CA INDEX NAME)



RN 184092-93-1 CAPLUS

CN Butanoic acid, 2-cyano-2-(dicyanomethyl)-4,4-diethoxy-, methyl ester (9CI) (CA INDEX NAME)



REFERENCE COUNT: 9 THERE ARE 9 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L6 ANSWER 20 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1996:664756 CAPLUS

DOCUMENT NUMBER: 125:329472

TITLE: Preparation of ring-fused pyrimidine-containing amino acid derivatives as antiprotozoan agents

INVENTOR(S): Horii, Toshihiro; Aono, Tetsuya

PATENT ASSIGNEE(S): Takeda Chemical Industries Ltd, Japan

SOURCE: Jpn. Kokai Tokkyo Koho, 31 pp.

CODEN: JKXXAF

DOCUMENT TYPE: Patent

LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 08225574	A	19960903	JP 1995-330939	19951220 <--
PRIORITY APPLN. INFO.:			JP 1995-330939	A 19951220
			JP 1994-317938	19941221

OTHER SOURCE(S): MARPAT 125:329472

GI For diagram(s), see printed CA Issue.

AB The title compds. [I; ring A = (un)substituted five membered ring; Z = (un)substituted bivalent aliphatic chain consisting of a series of <5 atoms and optionally interrupted by one hetero atom in the chain; B = (un)substituted 5- or 6-membered heterocyclyl or carbocyclyl; the substituent of B is preferably CONHCH(CO2R3)(CH2)pWR4; wherein p = 1-4; W = bond, O, NHCONH, NR, NRCO, CONR, NHSO2; wherein R = H, C1-4 hydrocarbyl; CO2R3 = optionally esterified CO2H; R4 = (un)substituted chain or cyclic group; or Z = (CR1R2)n-Z1; wherein R1, R2 = H, lower alkyl; Z1 = bond, O,

NH; n = 1-5] or salts thereof, which are useful for treating infections of protozoa, particularly coccidium and drug-resistant malaria, are prepared Thus, 4-[2-(2,4-diamino-7H-pyrrolo[2,3-d]pyrimidin-5-yl)ethyl]benzoic acid ditrifluoroacetate was condensed with Me O-(4-methoxycarbonylbenzyl)-L-serinate hydrochloride using di-Et cyanophosphate and Et3N in DMF at room temperature for 1 h to give 66% Me N-[4-[2-(2,4-diamino-7H-pyrrolo[2,3-d]pyrimidin-5-yl)ethyl]benzoyl]-O-O-(4-methoxycarbonylbenzyl)-L-serinate, which was saponified with a mixture of 1 h aqueous NaOH and MeOH at room temperature for

5 h and neutralized with dilute HCl to give 84% N-[4-[2-(2,4-diamino-7H-pyrrolo[2,3-d]pyrimidin-5-yl)ethyl]benzoyl]-O-O-(4-methoxycarbonylbenzyl)-L-serine. These compds. I inhibited dihydrofolic acid reductase of malaria protozoa Plasmodium falciparum with IC50 of 0.8-62 nM. The title compound (II; R5 = Q) showed ED50 of 0.17 and 0.11 nM for inhibiting the proliferation of wild type-malaria protozoa P. falciparum 3D7 and cycloguanyl-resistant P. falciparum FCR3, resp. Capsule, tablet, and vial formulations containing II (R5 = Q1) were prepared

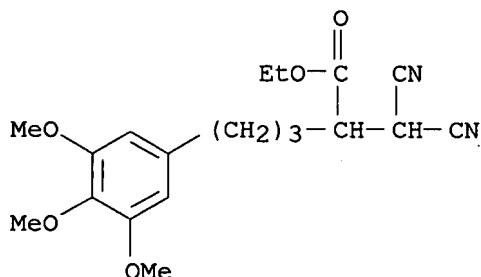
IT 182961-44-0P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

(preparation of ring-fused pyrimidine-containing amino acid derivs. as antiprotozoan agents)

RN 182961-44-0 CAPLUS

CN Benzenepentanoic acid, α -(dicyanomethyl)-3,4,5-trimethoxy-, ethyl ester (9CI) (CA INDEX NAME)



L6 ANSWER 21 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1996:365832 CAPLUS

DOCUMENT NUMBER: 125:86845

TITLE: Cyclopropenation and Related Reactions of Ruthenium Vinylidene Complexes

AUTHOR(S): Ting, Pei-Chen; Lin, Ying-Chih; Lee, Gene-Hsiang; Cheng, Ming-Chu; Wang, Yu

CORPORATE SOURCE: Department of Chemistry, National Taiwan University, Taipei, 106, Taiwan

SOURCE: Journal of the American Chemical Society (1996), 118(27), 6433-6444

CODEN: JACSAT; ISSN: 0002-7863

PUBLISHER: American Chemical Society

DOCUMENT TYPE: Journal

LANGUAGE: English

AB Facile deprotonation of a number of cationic ruthenium vinylidene complexes, followed by cyclopropenation, is accomplished in acetone. The deprotonation of $[\text{Ru}]:\text{C}:(\text{Ph})\text{CH}_2\text{R}^+$, ($[\text{Ru}] = (\eta^5\text{-C}_5\text{H}_5)(\text{PPh}_3)_2\text{Ru}$ through out this abstract) by $n\text{-Bu}_4\text{NOH}$ induces a novel cyclization reaction and yields the neutral cyclopropenyl complexes [cyclic] $[\text{Ru}]-\text{C}:\text{C}(\text{Ph})\text{CHR}$ (3b, $\text{R} = \text{CN}$; 3c, $\text{R} = \text{Ph}$; 3d, $\text{R} = \text{CH}:\text{CH}_2$; 3e, $\text{R} = \text{CH}:\text{CMe}_2$). Cyclic complex $[\text{Ru}]-\text{C}:\text{C}(\text{C}_6\text{H}_9)\text{CHCN}^+$ is similarly prepared Protonation of 3b-3e regenerates

the corresponding vinylidene complexes. Deprotonation of $[\text{Ru}]:\text{C}:\text{C}(\text{Ph})\text{CH}_2\text{COOMe}^+$ by $n\text{-Bu}_4\text{NOH}$ induces a different type of cyclization and yields the neutral furan complex [cyclic] $[\text{Ru}]-\text{C}:\text{C}(\text{Ph})\text{CH}:\text{C}(\text{O})\text{OMe}$ (4h). The cyclopropenyl complex containing a methoxy substituent cannot be prepared from $[\text{Ru}]:\text{C}:\text{C}(\text{Ph})\text{CH}_2\text{OCH}_3^+$ (2i), but F^- of $n\text{-Bu}_4\text{NF}$ attacks the C_α of 2i to produce the unstable vinyl complex $[\text{Ru}]\text{C}(\text{F}):\text{C}(\text{Ph})\text{CH}_2\text{OCH}_3$. Cyclic complex $[\text{Ru}]-\text{C}:\text{C}(\text{Ph})\text{C}(\text{CN})\text{OCH}_3$ (9b) was indirectly prepared from the addition of TCNQ to 3b, giving [cyclic] $[\text{Ru}]:\text{C}:\text{C}(\text{Ph})\text{CH}(\text{CN})\text{TCNQ}$ (6b) followed by methanolysis. Unlike 3, complex 9b is not converted to vinylidene complex, instead, removal of the methoxy substituent by acid gives the cationic cyclopropenyl complex $[\text{Ru}]-\text{C}:\text{C}(\text{Ph})\text{C}(\text{CN})^+$. Cyclic complex $[\text{Ru}]-\text{C}:\text{C}(\text{Ph})\text{C}(\text{COOMe})^+$ is similarly prepared from 4h via a TCNQ complex followed by a methoxy-substituted complex. In the presence of allyl iodide, opening of the three-membered ring of 3b, followed by a subsequent oxidative coupling reaction, gives a dimeric dicationic product $\{[\text{Ru}]:\text{C}:\text{C}(\text{Ph})-\text{CHCN}\}_2^{2+}$ (11). Proton abstraction of 11 by $n\text{-Bu}_4\text{NOH}$ gives the biscyclopropenyl complex $\{[\text{Ru}]-\text{C}:\text{C}(\text{Ph})\text{CCN}\}_2$. Mol. structures of complexes 3b, 4h, 6b, 9b, 11, and [cyclic] $[\text{Ru}]-\text{C}:\text{C}(\text{Ph})\text{C}(\text{CPh}_3)\text{CN}$ have been confirmed by x-ray diffraction anal.

IT 178687-62-2P

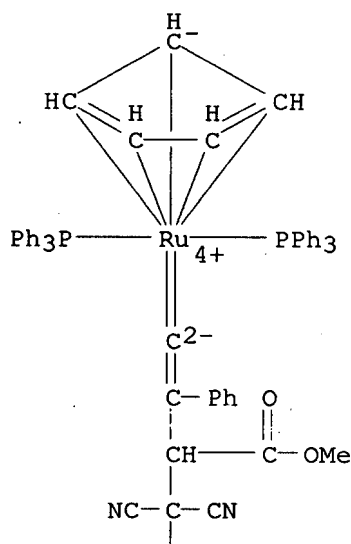
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

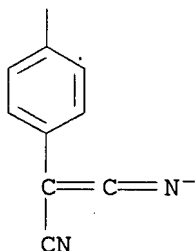
(cyclopropenation and related reactions of ruthenium vinylidene complexes)

RN 178687-62-2 CAPLUS

CN Ruthenium, $(\eta^5\text{-}2,4\text{-cyclopentadien-1-yl})[4,4\text{-dicyano-}4\text{-[4-(cyanoiminatoethenyl)phenyl]-3-(methoxycarbonyl)-2-phenyl-1-butenylidene]bis(triphenylphosphine)-$ (9CI) (CA INDEX NAME)

PAGE 1-A





L6 ANSWER 22 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1996:6671 CAPLUS

DOCUMENT NUMBER: 124:177091

TITLE: Novel gem-dinitrile functionalized polyesters and polyamides from malononitrile; potential piezoelectric materials

AUTHOR(S): Steadman, Scott; Parrish, Dennis A.; Mathias, Lon J.

CORPORATE SOURCE: Department of Polymer Science, University of Southern Mississippi, Hattiesburg, MS, 39406-0076, USA

SOURCE: Polymer Preprints (American Chemical Society, Division of Polymer Chemistry) (1995), 36(2), 320-1
CODEN: ACPPAY; ISSN: 0032-3934

PUBLISHER: American Chemical Society, Division of Polymer Chemistry

DOCUMENT TYPE: Journal

LANGUAGE: English

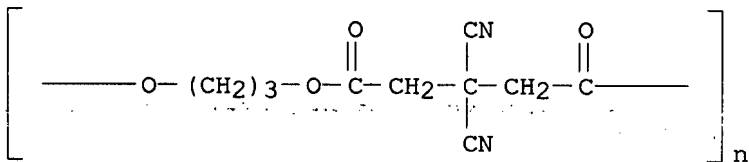
AB Novel polyamides and polyesters in which the dinitrile group can potentially align in the same direction as the dipole of the carbonyl groups were synthesized via step growth dialkylation of malononitrile. Diamide and diester monomers facilitated polymerization by the attachment of chlorine to an activated position (α to carbonyl). The polymers, having mol. weight 5000-8000, were characterized by NMR, viscosity and thermal anal.

IT 169893-85-0P 174297-80-4P 174297-82-6P

RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)
(preparation of gem-dinitrile functionalized polyesters and polyamides from malononitrile as potential piezoelec. materials)

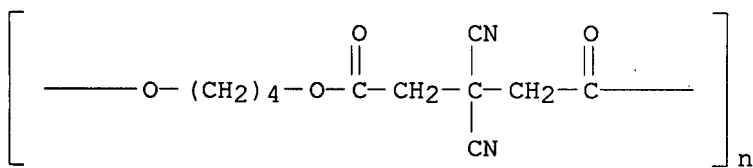
RN 169893-85-0 CAPLUS

CN Poly[oxy-1,3-propanediyl oxy(3,3-dicyano-1,5-dioxo-1,5-pentanediy)] (9CI)
(CA INDEX NAME)

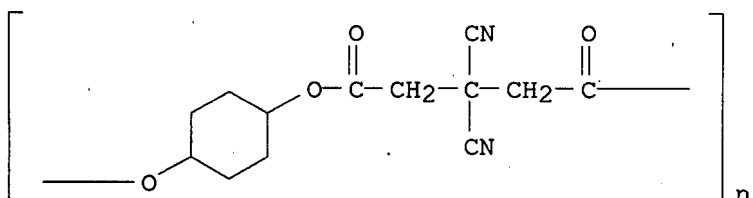


RN 174297-80-4 CAPLUS

CN Poly[oxy-1,4-butanediyl oxy(3,3-dicyano-1,5-dioxo-1,5-pentanediy)] (9CI)
(CA INDEX NAME)

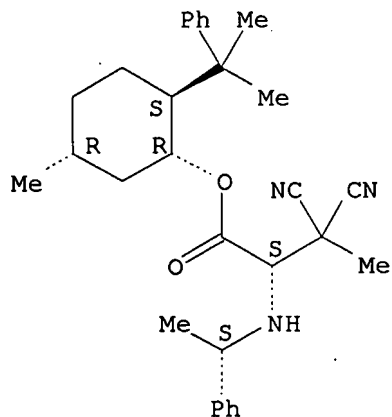


RN 174297-82-6 CAPLUS
 CN Poly[oxy-1,4-cyclohexanedioxy(3,3-dicyano-1,5-dioxo-1,5-pentanedioyl)]
 (9CI) (CA INDEX NAME)



L6 ANSWER 23 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN
 ACCESSION NUMBER: 1995:900662 CAPLUS
 DOCUMENT NUMBER: 124:116317
 TITLE: Lanthanum isopropoxide catalyzed addition of activated nucleophiles to imines
 AUTHOR(S): Yamamoto, Yoshinori; Fukui, Hiroyuki; Honda, Yoshihiro
 CORPORATE SOURCE: Dept. Chem., Tohoku Univ., Sendai, 980-77, Japan
 SOURCE: Applied Organometallic Chemistry (1995), 9(5 & 6), 467-71
 CODEN: AOCHEX; ISSN: 0268-2605
 PUBLISHER: Wiley
 DOCUMENT TYPE: Journal
 LANGUAGE: English
 OTHER SOURCE(S): CASREACT 124:116317
 AB The addition of certain activated nucleophiles to activated imines is catalyzed by lanthanum isopropoxide. As activated nucleophiles, methylmalonitrile and Me 2-cyanopropanoate can be utilized. Imines having an electron-withdrawing group either at the carbon or at the nitrogen atom of the C:N double bond can be used: for example N-toluenesulfonylimines, N-(4-methoxycarbonylphenyl)imines and α-imino esters.
 IT 155751-02-3P
 RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)
 (lanthanum isopropoxide catalyzed addition of activated nucleophiles to imines)
 RN 155751-02-3 CAPLUS
 CN Butanoic acid, 3,3-dicyano-2-[(1-phenylethyl)amino]--, 5-methyl-2-(1-methyl-1-phenylethyl)cyclohexyl ester, [1R-[1α[S*(S*)],2β,5α]]-
 (9CI) (CA INDEX NAME)

Absolute stereochemistry.



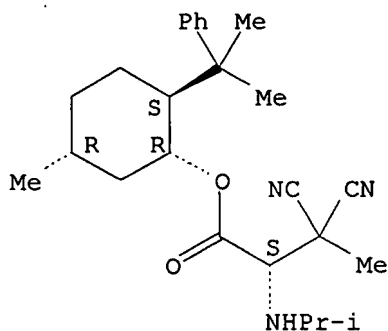
IT 155696-71-2P 155696-72-3P 172880-55-6P
 172880-56-7P 173006-24-1P 173006-25-2P
 173006-26-3P

RL: SPN (Synthetic preparation); PREP (Preparation)
 (lanthanum isopropoxide catalyzed addition of activated nucleophiles to
 imines)

RN 155696-71-2 CAPLUS

CN Butanoic acid, 3,3-dicyano-2-[(1-methylethyl)amino]-, 5-methyl-2-(1-methyl-
 1-phenylethyl)cyclohexyl ester, [1R-[1 α (S*),2 β ,5 α]]-
 (9CI) (CA INDEX NAME)

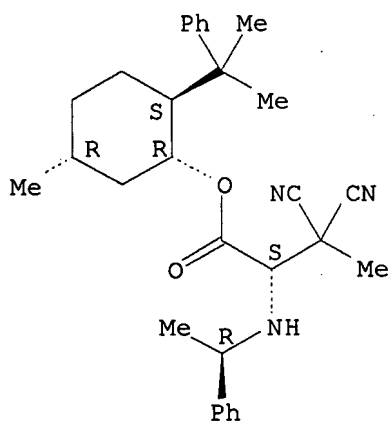
Absolute stereochemistry.



RN 155696-72-3 CAPLUS

CN Butanoic acid, 3,3-dicyano-2-[(1-phenylethyl)amino]-, 5-methyl-2-(1-methyl-
 1-phenylethyl)cyclohexyl ester, [1R-[1 α [S*(R*)],2 β ,5 α]]-
 (9CI) (CA INDEX NAME)

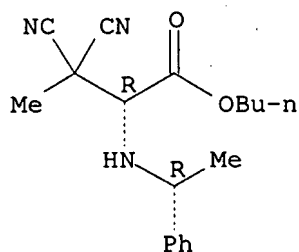
Absolute stereochemistry.



RN 172880-55-6 CAPLUS

CN Butanoic acid, 3,3-dicyano-2-[(1-phenylethyl)amino]-, butyl ester,
(R*,R*)- (9CI) (CA INDEX NAME)

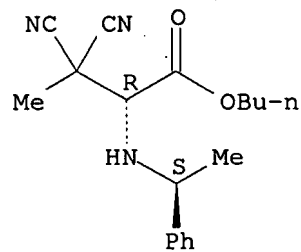
Relative stereochemistry.



RN 172880-56-7 CAPLUS

CN Butanoic acid, 3,3-dicyano-2-[(1-phenylethyl)amino]-, butyl ester,
(R*,S*)- (9CI) (CA INDEX NAME)

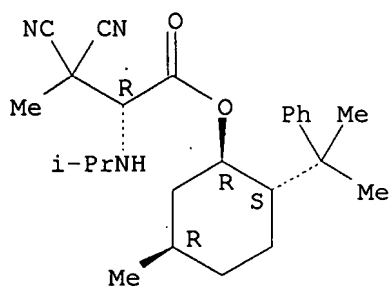
Relative stereochemistry.



RN 173006-24-1 CAPLUS

CN Butanoic acid, 3,3-dicyano-2-[(1-methylethyl)amino]-, 5-methyl-2-(1-methyl-1-phenylethyl)cyclohexyl ester, [1R-[1α(R*),2β,5α]]-
(9CI) (CA INDEX NAME)

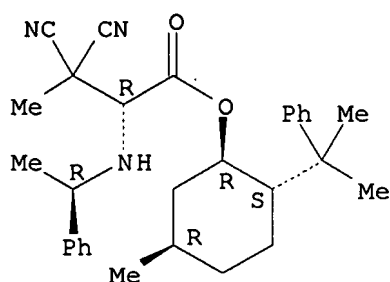
Absolute stereochemistry.



RN 173006-25-2 CAPLUS

CN Butanoic acid, 3,3-dicyano-2-[(1-phenylethyl)amino]-, 5-methyl-2-(1-methyl-1-phenylethyl)cyclohexyl ester, [1R-[1α[R*(R*)],2β,5α]]-
(9CI) (CA INDEX NAME)

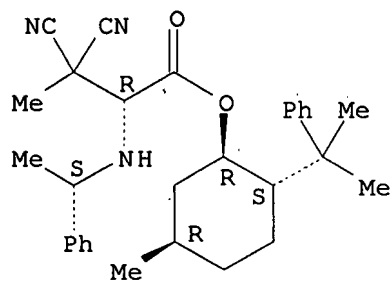
Absolute stereochemistry.



RN 173006-26-3 CAPLUS

CN Butanoic acid, 3,3-dicyano-2-[(1-phenylethyl)amino]-, 5-methyl-2-(1-methyl-1-phenylethyl)cyclohexyl ester, [1R-[1α[R*(S*)],2β,5α]]-
(9CI) (CA INDEX NAME)

Absolute stereochemistry.



L6 ANSWER 24 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1995:845626 CAPLUS

DOCUMENT NUMBER: 124:86769

TITLE: Novel three-component reaction of 1,1-dicyano-2-(trifluoromethyl)ethylenes with primary arylamines and ketones

AUTHOR(S): Tyutin, V. Yu.; Chkanikov, N. D.; Nesterov, V. N.; Antipin, M. Yu.; Struchkov, Yu. T.; Kolomiets, A. F.; Fokin, A. V.

CORPORATE SOURCE: A. N. Nesmeyanov Inst. Organoelem. Compd., Russ. Acad. Sci., Moscow, 117813, Russia

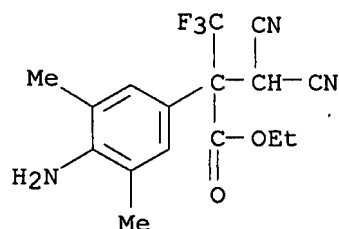
SOURCE: Izvestiya Akademii Nauk, Seriya Khimicheskaya (1993), (3), 552-9
 CODEN: IASKEA
 PUBLISHER: Nauka
 DOCUMENT TYPE: Journal
 LANGUAGE: Russian
 OTHER SOURCE(S): CASREACT 124:86769

AB 1,1-Dicyano-2,2-bis(trifluoromethyl)ethylene and 3,3-dicyano-2-(trifluoromethyl)acrylates react with primary arylamines in the presence of ketones to form 1,1-aryl-1,4-dihydropyridine derivs. under mild conditions. In this three-component reaction Schiff bases are formed as intermediates. 1,4-Dihydropyridines derivs., which are the products of three-component heterocyclization, were also obtained by interaction of the corresponding Schiff bases with 1,1-dicyano-2-(trifluoromethyl)ethylenes.

IT 134641-39-7P
 RL: SPN (Synthetic preparation); PREP (Preparation)
 (preparation of)

RN 134641-39-7 CAPLUS

CN Benzeneacetic acid, 4-amino- α -(dicyanomethyl)-3,5-dimethyl- α -(trifluoromethyl)-, ethyl ester (9CI) (CA INDEX NAME)



L6 ANSWER 25 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1995:695727 CAPLUS

DOCUMENT NUMBER: 123:286966

TITLE: Novel gem-dinitrile functionalized polyesters and polyamides from malononitrile; potential piezoelectric materials

AUTHOR(S): Mathias, Lon J.; Parrish, Dennis A.; Steadman, Scott
 CORPORATE SOURCE: Department Polymer Science, University Southern Mississippi, Hattiesburg, MS, 39406-0076, USA

SOURCE: Polymer Preprints (American Chemical Society, Division of Polymer Chemistry) (1994), 35(2), 659-60
 CODEN: ACPPAY; ISSN: 0032-3934

PUBLISHER: American Chemical Society, Division of Polymer Chemistry

DOCUMENT TYPE: Journal

LANGUAGE: English

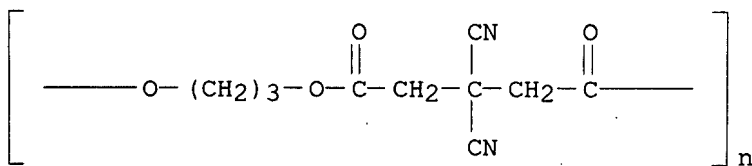
AB The initial success is described in obtaining a polyester and polyamide in which the dinitrile group net dipole can potentially align in the same direction as the carbonyl groups.

IT 169893-85-0P

RL: SPN (Synthetic preparation); PREP (Preparation)
 (preparation of gem-dinitrile functionalized polyesters and polyamides from malononitrile as potential piezoelec. materials)

RN 169893-85-0 CAPLUS

CN Poly[oxy-1,3-propanediyl-oxy(3,3-dicyano-1,5-dioxo-1,5-pentanediy)] (9CI)
 (CA INDEX NAME)



L6 ANSWER 26 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1995:440268 CAPLUS

DOCUMENT NUMBER: 123:112653

TITLE: Synthesis and antitumor activity of pyrrolo[2,3-d]pyrimidine antifolates with a bridge chain containing a nitrogen atom

AUTHOR(S): Aso, Kazuyoshi; Hitaka, Takenori; Yukishige, Koichi; Ootsu, Koichiro; Akimoto, Hiroshi

CORPORATE SOURCE: Pharmaceutical Res. Div., Takeda Chem. Industries, Ltd., Osaka, 532, Japan

SOURCE: Chemical & Pharmaceutical Bulletin (1995), 43(2), 256-61

CODEN: CPBTAL; ISSN: 0009-2363

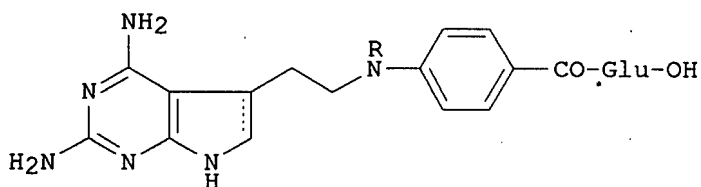
PUBLISHER: Pharmaceutical Society of Japan

DOCUMENT TYPE: Journal

LANGUAGE: English

OTHER SOURCE(S): CASREACT 123:112653

GI



I

AB Novel pyrrolo[2,3-d]pyrimidine antifolates I (R = H, Me) with a nitrogen atom in the bridge chain between the 2,4-diaminopyrrolo[2,3-d]pyrimidine and phenylene rings were designed and efficiently synthesized. I exhibited more potent inhibitory activities than methotrexate (MTX) against the proliferation of human epidermoid carcinoma KB cells and human non-small cell lung carcinoma A549 cells despite their modest dihydrofolate reductase (DHFR)-inhibitory potency.

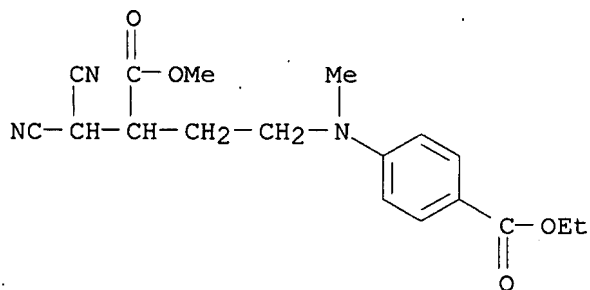
IT 133719-38-7P 133719-41-2P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)

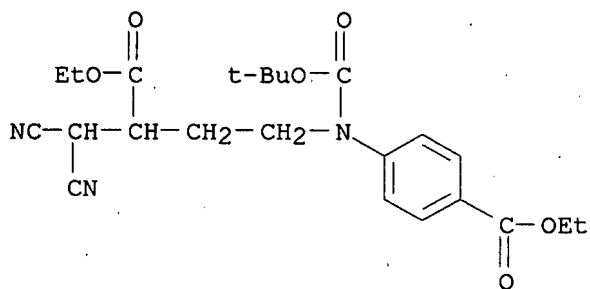
(preparation and antitumor activity of pyrrolo[2,3-d]pyrimidine antifolates with nitrogen-containing bridge chains)

RN 133719-38-7 CAPLUS

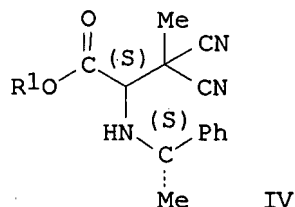
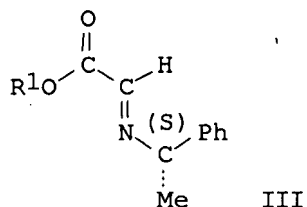
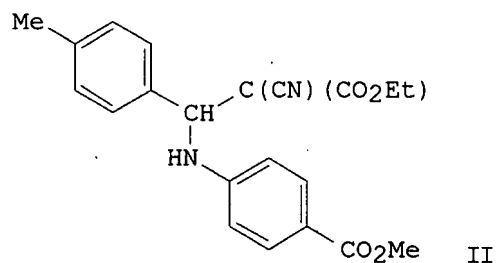
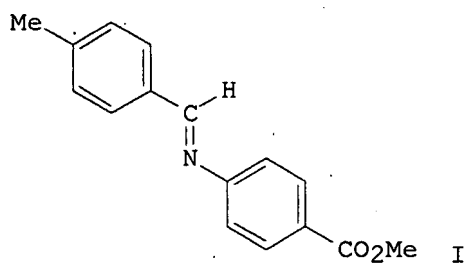
CN Benzoic acid, 4-[[4,4-dicyano-3-(methoxycarbonyl)butyl]methylanino]-, ethyl ester (9CI) (CA INDEX NAME)



RN 133719-41-2 CAPLUS
 CN Benzoic acid, 4-[[4,4-dicyano-3-(ethoxycarbonyl)butyl][(1,1-dimethylethoxy)carbonyl]amino]-, ethyl ester (9CI) (CA INDEX NAME)



L6 ANSWER 27 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN
 ACCESSION NUMBER: 1994:434408 CAPLUS
 DOCUMENT NUMBER: 121:34408
 TITLE: Transition metal catalyzed addition of certain nucleophiles to imines
 AUTHOR(S): Yamamoto, Yoshinori; Kubota, Yasufumi; Honda, Yoshihiro; Fukui, Hiroyuki; Asao, Naoki; Nemoto, Hisao
 CORPORATE SOURCE: Faculty of Science, Tohoku University, Sendai, 980, Japan
 SOURCE: Journal of the American Chemical Society (1994), 116(7), 3161-2
 CODEN: JACSAT; ISSN: 0002-7863
 DOCUMENT TYPE: Journal
 LANGUAGE: English
 OTHER SOURCE(S): CASREACT 121:34408
 GI



AB Imines react with certain nucleophiles in the presence of catalytic amts. of transition metal complexes to give alkylation products in good yield. Thus, imine (I) was treated with $\text{CH}(\text{CN})_2(\text{CO}_2\text{Et})$ in the presence of $\text{RhHCO}(\text{PPh}_3)_3$ in THF solvent to give alkylation product II in 75% yield. A significantly high diastereomeric excess was accomplished by using III [R1 = (-)-8-phenylmenthyl] in which a chiral auxiliary exists at the ester unit. The $\text{Ls}(\text{O-iso-Pr})_3$ catalyzed reaction of III with $\text{CH}(\text{CN})_2\text{Me}$ in THF at room temperature gave IV (R1 as above) as the predominant diastereoisomer in a 90:10 ratio; x-ray anal. indicate that the α -carbon to the amino group possesses the S configuration.

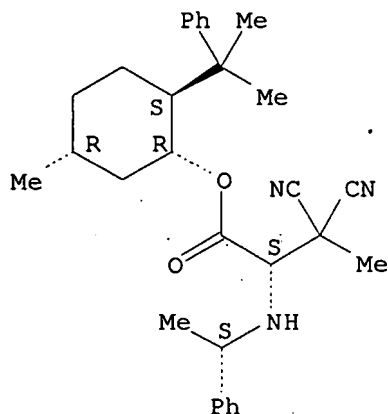
IT 155751-02-3P

RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of, absolute configuration of)

RN 155751-02-3 CAPLUS

CN Butanoic acid, 3,3-dicyano-2-[(1-phenylethyl)amino]-, 5-methyl-2-(1-methyl-1-phenylethyl)cyclohexyl ester, [1R-[1 α [S*(S*)],2 β ,5 α]]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



IT 155696-71-2P 155696-72-3P

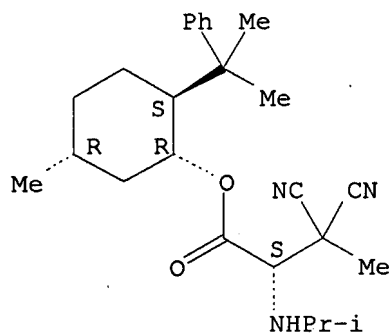
RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of, stereoselective)

RN 155696-71-2 CAPLUS

CN Butanoic acid, 3,3-dicyano-2-[(1-methylethyl)amino]-, 5-methyl-2-(1-methyl-

1-phenylethyl)cyclohexyl ester, [1R-[1 α (S*),2 β ,5 α]]-(9CI) (CA INDEX NAME)

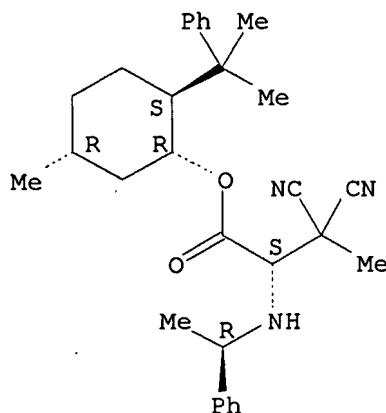
Absolute stereochemistry.



RN 155696-72-3 CAPLUS

CN Butanoic acid, 3,3-dicyano-2-[(1-phenylethyl)amino]-, 5-methyl-2-(1-methyl-1-phenylethyl)cyclohexyl ester, [1R-[1 α [S*(R*)],2 β ,5 α]]-(9CI) (CA INDEX NAME)

Absolute stereochemistry.



L6 ANSWER 28 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1994:269808 CAPLUS

DOCUMENT NUMBER: 120:269808

TITLE: Wittig reactions of a fluoren-9-ylidene and an anthrone-10-arylidene

AUTHOR(S): Ganoub, Neven A. F.

CORPORATE SOURCE: Dep. Pesticide Chem., Natl. Res. Cent., Cairo, Egypt

SOURCE: Phosphorus, Sulfur and Silicon and the Related Elements (1993), 81(1-4), 125-31

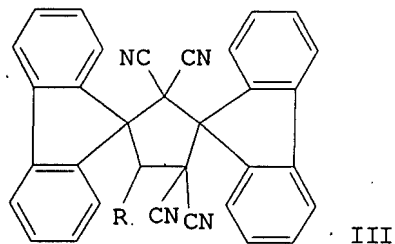
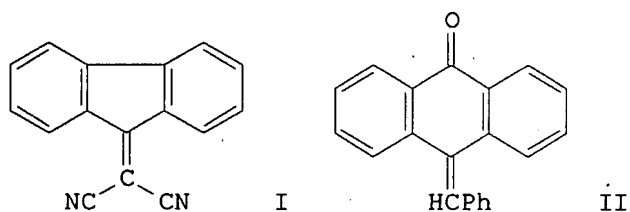
CODEN: PSSLEC; ISSN: 1042-6507

DOCUMENT TYPE: Journal

LANGUAGE: English

OTHER SOURCE(S): CASREACT 120:269808

GI

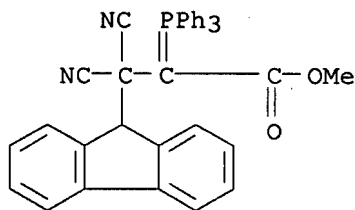


AB The Wittig reactions of fluorene-9-ylidenemalonitrile (I) and 10-benzylideneanthrone (II) with phosphonium ylides $\text{Ph}_3\text{P}+\text{CH}-\text{CO}_2\text{R}$ ($\text{R} = \text{Me}, \text{Et}$) have been investigated. In both cases, unusual reaction products, e.g., bis(9-fluorenyl)cyclopentane III (from I), were isolated and identified on the basis of elemental analyses and spectral studies.

IT 154496-99-8P 154497-00-4P
 RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
 (preparation and thermal decomposition of)

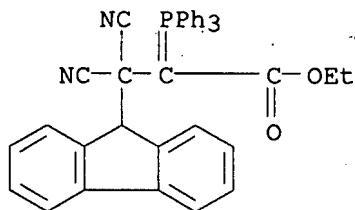
RN 154496-99-8 CAPLUS

CN 9H-Fluorene-9-propanoic acid, β,β -dicyano- α -(triphenylphosphoranylidene)-, methyl ester (9CI) (CA INDEX NAME)



RN 154497-00-4 CAPLUS

CN 9H-Fluorene-9-propanoic acid, β,β -dicyano- α -(triphenylphosphoranylidene)-, ethyl ester (9CI) (CA INDEX NAME)



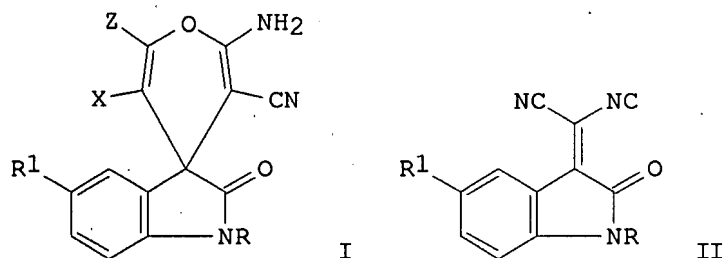
L6 ANSWER 29 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1994:244629 CAPLUS

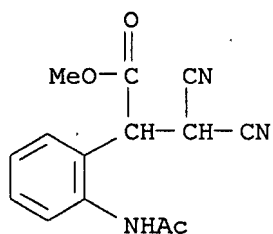
DOCUMENT NUMBER: 120:244629

TITLE: Synthesis of spiro indolin-2-one derivatives

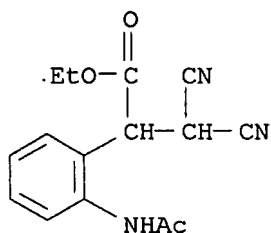
AUTHOR(S): El-Ahl, Abdel Aziz S.; Afeefy, Hussein; Metwally, Mohamed Abbas
 CORPORATE SOURCE: Fac. Sci., Mansoura Univ., Mansoura, Egypt
 SOURCE: Journal of Chemical Research, Synopses (1994), (1), 14-15
 CODEN: JRPSDC; ISSN: 0308-2342
 DOCUMENT TYPE: Journal
 LANGUAGE: English
 OTHER SOURCE(S): CASREACT 120:244629
 GI



AB Title compds. I were prepared by heating dicyanomethyleneindolinones II (R = H, R1 = H, Me; R = Me, Ac, R1 = H) with active methylene compds., XCH2COZ (X = Ac, Z = OEt, Me; X = cyano, Z = Ph).
 IT 154379-70-1P 154379-71-2P
 RL: SPN (Synthetic preparation); PREP (Preparation)
 (preparation of)
 RN 154379-70-1 CAPLUS
 CN Benzeneacetic acid, 2-(acetylamino)- α -(dicyanomethyl)-, methyl ester (9CI) (CA INDEX NAME)



RN 154379-71-2 CAPLUS
 CN Benzeneacetic acid, 2-(acetylamino)- α -(dicyanomethyl)-, ethyl ester (9CI) (CA INDEX NAME)



L6 ANSWER 30 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1993:213100 CAPLUS
DOCUMENT NUMBER: 118:213100
TITLE: Preparation of tricyclic fused pyrimidine compounds
INVENTOR(S): Akimoto, Hiroshi; Otsu, Koichiro; Miwa, Tetsuo
PATENT ASSIGNEE(S): Takeda Chemical Industries, Ltd., Japan
SOURCE: Jpn. Kokai Tokkyo Koho, 25 pp.
CODEN: JKXXAF
DOCUMENT TYPE: Patent
LANGUAGE: Japanese
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 04211063	A	19920803	JP 1991-65613	19910305 <--
PRIORITY APPLN. INFO.:			JP 1990-54620	A1 19900305
OTHER SOURCE(S):	MARPAT 118:213100			

GI For diagram(s), see printed CA Issue.

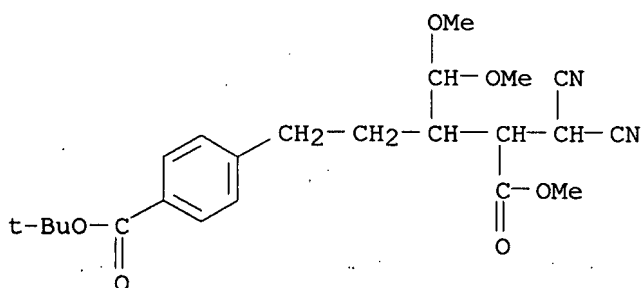
AB The title compds. [I; Q1 = H, halo, radical linked through C, N, O, or S; one of Q2 and Q3 = N, the other = N, CH; Y = N, CR1 (wherein R1 = H, hydrocarbyl), methylidyne; Z = C2-5 bivalent radical containing optional substituents; ring A1, A2 = (substituted) 5-7-membered ring; B = (substituted) cyclic radical, etc.], useful as antitumor agents with high selectivity, are prepared Cyclocondensation of 1.181 g ester II (preparation given) with 314 mg guanidine HCl and Me3COK in Me3COH gave 1.02 g pyrrolopyrimidine III, which (1.010 g) was treated with borane-THF complex in THF at 0° and then at 50°, the solution cooled and stirred with HOAc-MeOH at room temperature to give 542 mg IV. The preferred doses of I are 2.0-500 mg/kg-day orally and 1.0-200 mg/kg injection.

IT 147239-87-0P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation and reaction of, in preparation of antitumor agent)

RN 147239-87-0 CAPLUS

CN Benzenepentanoic acid, α -(dicyanomethyl)- β -(dimethoxymethyl)-4-[(1,1-dimethylethoxy)carbonyl]-, methyl ester (9CI) (CA INDEX NAME)

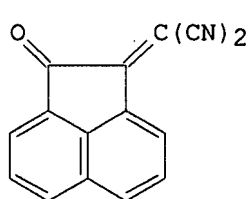


L6 ANSWER 31 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN

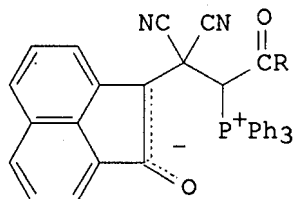
ACCESSION NUMBER: 1993:124655 CAPLUS
DOCUMENT NUMBER: 118:124655
TITLE: Wittig reaction of 1-(dicyanomethylene)acenaphthen-2-one
AUTHOR(S): Abdou, Wafaa M.; Ganoub, Neven A. F.
CORPORATE SOURCE: Natl. Res. Cent., Cairo, Egypt
SOURCE: Heteroatom Chemistry (1992), 3(2), 133-7
CODEN: HETCE8; ISSN: 1042-7163

DOCUMENT TYPE:
 LANGUAGE:
 OTHER SOURCE(S):
 GI

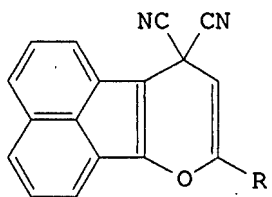
Journal
 English
 CASREACT 118:124655



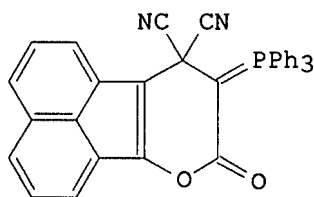
I



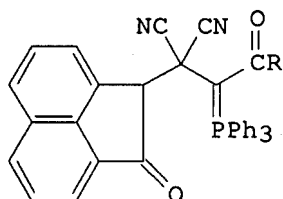
II



III



IV



V

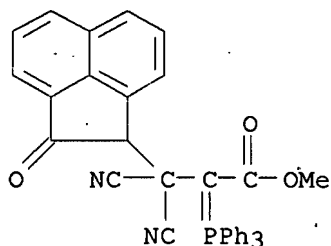
AB The Wittig reactions of title compound I with alkoxy carbonylmethylenetriphenylphosphoranes $\text{Ph}_3\text{P}+\text{C}=\text{HCOR}$ ($\text{R} = \text{OMe}, \text{OEt}$) were investigated and the reaction products zwitterion II and heterocycles III and IV were isolated. Reaction of I with benzoylmethylenetriphenylphosphorane $\text{Ph}_3\text{P}+\text{C}=\text{HCOPh}$ proceeded only at high temperature, yielding V and III ($\text{R} = \text{Ph}$). Mechanisms accounting for the formation of the adducts are discussed. Wittig olefination of several products was studied.

IT 145882-80-0P 145882-83-3P

RL: SPN (Synthetic preparation); PREP (Preparation)
 (preparation of)

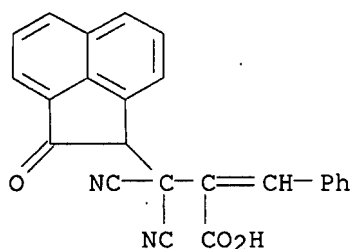
RN 145882-80-0 CAPLUS

CN 1-Acenaphthylenepropanoic acid, β,β -dicyano-1,2-dihydro-2-oxo- α -(triphenylphosphoranylidene)-, methyl ester (9CI) (CA INDEX NAME)

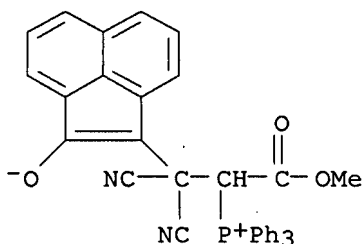


RN 145882-83-3 CAPLUS

CN 1-Acenaphthylenepropanoic acid, β,β -dicyano-1,2-dihydro-2-oxo- α -(phenylmethylene)- (9CI) (CA INDEX NAME)



IT 145882-82-2P
 RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
 (preparation, Wittig olefination, and thermal intramol. cyclocondensation of)
 RN 145882-82-2 CAPLUS
 CN Phosphonium, [2,2-dicyano-2-(2-hydroxy-1-acenaphthylenyl)-1-(methoxycarbonyl)ethyl]triphenyl-, inner salt (9CI) (CA INDEX NAME)



L6 ANSWER 32 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN
 ACCESSION NUMBER: 1993:29863 CAPLUS
 DOCUMENT NUMBER: 118:29863
 TITLE: Silver halide photographic material containing a compound which releases photographically useful species upon development
 INVENTOR(S): Asatake, Atsushi
 PATENT ASSIGNEE(S): Konica Co., Japan
 SOURCE: Jpn. Kokai Tokkyo Koho, 19 pp.
 CODEN: JKXXAF
 DOCUMENT TYPE: Patent
 LANGUAGE: Japanese
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 04177243	A	19920624	JP 1990-305540	19901110 <--
PRIORITY APPLN. INFO.:			JP 1990-305540	19901110

AB The photog. material contains a compound CRR1R2C(R3R4)mZ(Z1)nPUG (R = leaving group released by nucleophilic substitution; R1,R2,R3,R4 = H, aliphatic, aromatic, heterocyclic or electron-attracting group; Z = electron-attracting group; Z1 = timing group to be subjected to break and release PUG = photog. useful group; m,n = 0, 1). The photog. material has good storage stability, while upon development, it releases the PUGs at a proper reaction rate even in developer solution of relatively low pH.

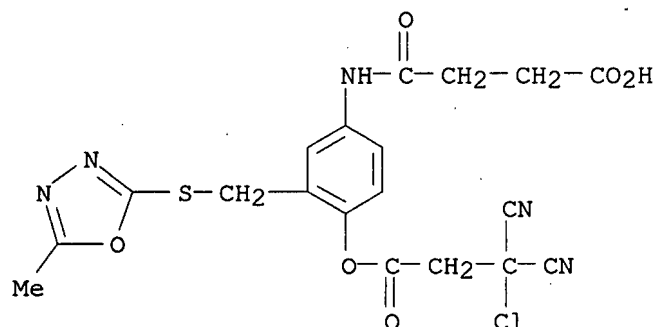
IT 144896-71-9 145059-42-3

RL: USES (Uses)

(photog. useful group-releasing, in processing)

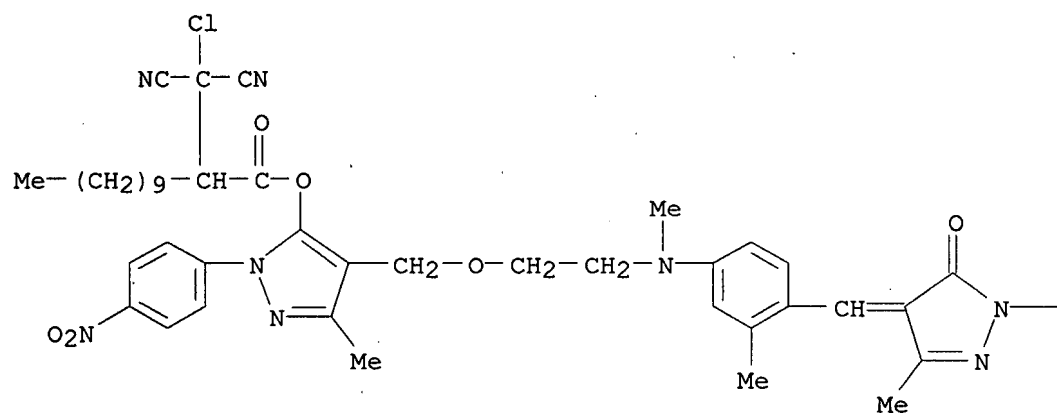
RN 144896-71-9 CAPLUS

CN Butanoic acid, 4-[[4-(3-chloro-3,3-dicyano-1-oxopropoxy)-3-[[5-methyl-1,3,4-oxadiazol-2-yl]thio]methyl]phenyl]amino]-4-oxo- (9CI) (CA INDEX NAME)



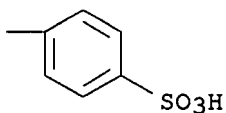
RN 145059-42-3 CAPLUS

CN Dodecanoic acid, 2-(chlorodicyanomethyl)-, 4-[[2-[[4-[[1,5-dihydro-3-methyl-5-oxo-1-(4-sulfohenyl)-4H-pyrazol-4-ylidene]methyl]-3-methylphenyl]methylamino]ethoxy]methyl]-3-methyl-1-(4-nitrophenyl)-1H-pyrazol-5-yl ester, monopotassium salt (9CI) (CA INDEX NAME)

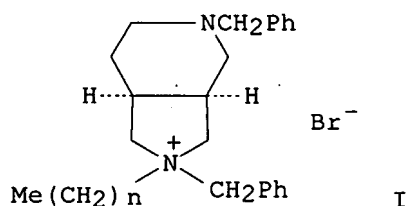


PAGE 1-A

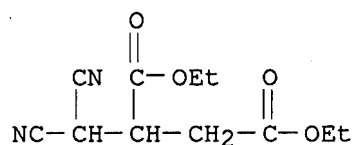
● K



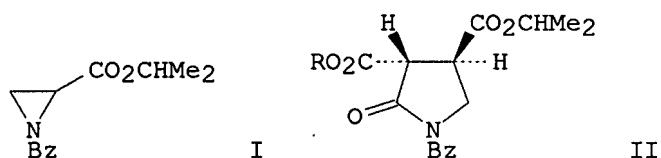
L6 ANSWER 33 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN
 ACCESSION NUMBER: 1992:174018 CAPLUS
 DOCUMENT NUMBER: 116:174018
 TITLE: Synthesis and structure-antimicrobial activity relationships of quaternary ammonium derivatives of perhydropyrrolo-[3,4-c]pyridine
 AUTHOR(S): Altomare, C.; Carotti, A.; Casini, G.; Cellamare, S.; Ferappi, M.; Vitali, C.
 CORPORATE SOURCE: Dip. Farm. Chim., Univ. Bari, Bari, I-70125, Italy
 SOURCE: Arzneimittel-Forschung (1992), 42(2), 152-5
 CODEN: ARZNAD; ISSN: 0004-4172
 DOCUMENT TYPE: Journal
 LANGUAGE: English
 OTHER SOURCE(S): CASREACT 116:174018
 GI



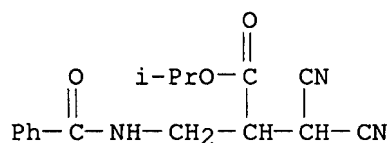
AB A homologous series of perhydropyrrolo[3,4-c]pyridine quaternary ammonium derivs. I ($n = 5, 7, 9-13, 15$) was synthesized from $\text{EtO}_2\text{CCH}_2\text{CH}(\text{CO}_2\text{Et})\text{CH}(\text{CN})_2$ and tested for in vitro antibacterial activity against different gram-pos. and gram-neg. bacteria. All I were more potent than the reference compound, benzalkonium chloride. Antibacterial activity, expressed as $\log 1/\text{MIC}$, was linearly related to lipophilicity up to C13-C14 homologs, where a break in the linear relationship was observed
 IT 82584-86-9
 RL: RCT (Reactant); RACT (Reactant or reagent)
 (hydrogenation-intramol. cyclocondensation of)
 RN 82584-86-9 CAPLUS
 CN Butanedioic acid, (dicyanomethyl)-, diethyl ester (9CI) (CA INDEX NAME)



L6 ANSWER 34 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN
 ACCESSION NUMBER: 1992:83465 CAPLUS
 DOCUMENT NUMBER: 116:83465
 TITLE: The regioselectivity of the ring opening of
 1-activated or nonactivated 2-alkoxycarbonyl or
 2-cyanoaziridines by carbanions of the dicarbonyl
 compounds.
 AUTHOR(S): Bouayad, Zoheir; Chanet-Ray, Josette; Ducher, S.;
 Vessiere, Roger
 CORPORATE SOURCE: Ec. Natl. Super. Chim. Clermont-Ferrand, Univ. Blaise
 Pascal, Aubiere, 63177, Fr.
 SOURCE: Journal of Heterocyclic Chemistry (1991),
 28(7), 1757-67
 CODEN: JHTCAD; ISSN: 0022-152X
 DOCUMENT TYPE: Journal
 LANGUAGE: English
 GI

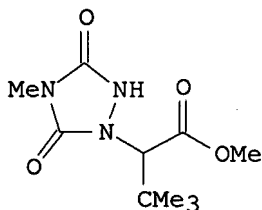


AB Aziridines, e.g. I, reacted with carbanions of dicarbonyl compds., e.g.
 $\text{RO}_2\text{CCH}_2\text{CO}_2\text{R}$ ($\text{R} = \text{Me}, \text{Et}, \text{CHMe}_2$), to give ring opened products and/or ring
 enlarged products, e.g. $(\text{RO}_2\text{C})_2\text{CHCH}_2\text{CH}(\text{NHBz})\text{CO}_2\text{CHMe}_2$,
 $(\text{RO}_2\text{C})_2\text{CHCH}(\text{CO}_2\text{CHMe}_2)\text{CH}_2\text{NHBz}$, and pyrrole II. The regioselectivity
 depends on several factors. The Ph group on C-3 favors C-3-N bond
 cleavage, whereas C-2-N bond cleavage is predominant with C-3 substituted
 or C-2-H aziridines. Cyanoaziridines are predominantly cleaved at C-3-N.
 IT 138478-35-0P
 RL: SPN (Synthetic preparation); PREP (Preparation)
 (preparation of)
 RN 138478-35-0 CAPLUS
 CN Propanoic acid, 2-[(benzoylamino)methyl]-3,3-dicyano-, 1-methylethyl ester
 (9CI) (CA INDEX NAME)



L6 ANSWER 35 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN
 ACCESSION NUMBER: 1991:680104 CAPLUS

DOCUMENT NUMBER: 115:280104
 TITLE: E/Z isomerization, solvolysis, addition, and cycloaddition reactions of (E)-tert-butylketene methyl tert-butyldimethylsilyl acetal
 AUTHOR(S): Adam, Waldemar; Wang, Xiaoheng
 CORPORATE SOURCE: Inst. Org. Chem., Univ. Wuerzburg, Wuerzburg, D-8700, Germany
 SOURCE: Journal of Organic Chemistry (1991), 56(26), 7244-50
 CODEN: JOCEAH; ISSN: 0022-3263
 DOCUMENT TYPE: Journal
 LANGUAGE: English
 OTHER SOURCE(S): CASREACT 115:280104
 GI



II

AB In the presence of catalytic amts. of CF₃COMe or CF₃COCF₃, the silyl ketene acetal Me₃CCCH:C(OMe)OSiMe₂CMe₃ (E-I) was isomerized into its Z isomer (Z/E ratio 90:10). For this novel E/Z isomerization a mechanism is proposed in which addition and reelimination of the fluoro ketone through a 1,4-dipolar intermediate operates. With the protic nucleophiles MeOH, CF₃CH₂OH, or PhOH, the ketene acetal E-I afforded the ortho esters Me₃CCCH₂C(OMe)(OR)OSiMe₂CMe₃ (R = Me, CF₃CH₂, Ph) as addition products, while AcOH, CF₃CO₂H, or H₂O led to Me pivalate as the solvolysis product. This chemical is readily explained through protonation of the ketene acetal E-I to generate the corresponding carbenium ion. At low temperature the reaction with TCNE gave the silylketene imine as labile cycloadduct, which underwent desilylation on workup to give the TCNE-incorporated ester (NC)₂CHC(CN)₂CH(CMe₂)C(O)OMe; the latter eliminated hydrogen cyanide at room temperature to give the ene ester. With MTAD the labile silyl ene product was obtained initially, which underwent silyl migration to give N-silylated urazole; final desilylation led to the stable urazole II. Also, for the ene reactions of TCNE and MTAD with the silyl ketene acetal E-I, intervention of a 1,4-dipolar intermediate is proposed.

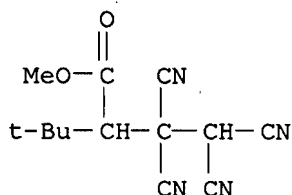
IT 136911-64-3P

RL: SPN (Synthetic preparation); PREP (Preparation)

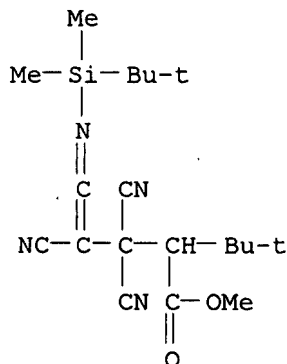
(preparation and hydrogen cyanide elimination of)

RN 136911-64-3 CAPLUS

CN Butanoic acid, 3,3,4,4-tetracyano-2-(1,1-dimethylethyl)-, methyl ester (9CI) (CA INDEX NAME)



IT 136911-63-2P
 RL: SPN (Synthetic preparation); PREP (Preparation)
 (preparation of)
 RN 136911-63-2 CAPLUS
 CN 4-Pentenoic acid, 3,3,4-tricyano-2-(1,1-dimethylethyl)-5-[[(1,1-dimethylethyl)dimethylsilyl]imino]-, methyl ester (9CI) (CA INDEX NAME)



L6 ANSWER 36 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1991:608100 CAPLUS

DOCUMENT NUMBER: 115:208100

TITLE: Chemistry of phosphorus ylides. 10. Reaction with phosphacumulenes. IV. Synthesis of pyran, phosphoranylidene, oxaphosphorin and oxazaphosphorin from the reaction of 1,3-dioxo- Δ^2, α -indanmalononitrile with phosphoranes and iminophosphoranes

AUTHOR(S): Soliman, Fouad M.; Said, Medhat M.

CORPORATE SOURCE: Natl. Res. Cent., Cairo, Egypt

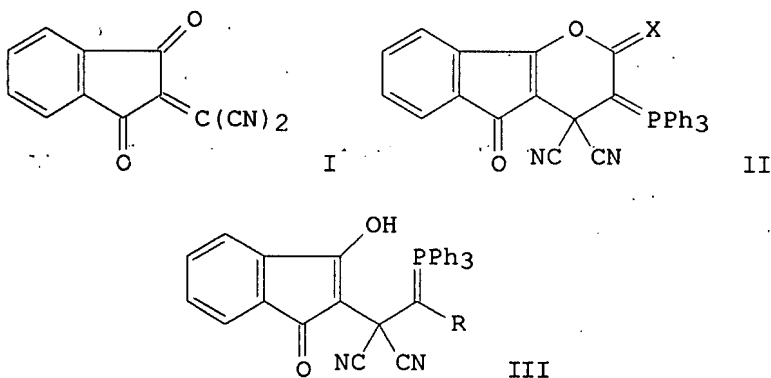
SOURCE: Phosphorus, Sulfur and Silicon and the Related Elements (1991), 61(3-4), 335-40
 CODEN: PSSLEC; ISSN: 1042-6507

DOCUMENT TYPE: Journal

LANGUAGE: English

OTHER SOURCE(S): CASREACT 115:208100

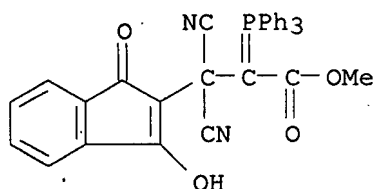
GI



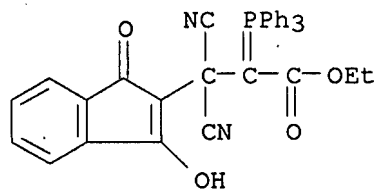
AB 1,3-Dioxo- Δ^2, α -indanmalononitrile (I) reacts with the active ketenylidene- and thioketenylidenetriphenylphosphoranes $\text{Ph}_3\text{P}:\text{C}:\text{C}:\text{X}$ ($\text{X} = \text{O},$

S, resp.) to give the corresponding pyrans II (X = O, S). The reaction of II with 4-O₂NC₆H₄CHO proceeds according to the Wittig reaction to give the resp. methylenide derivs. On the other hand, phosphoranylidenes III (R = acyl, alkoxy-carbonyl) were isolated from the reaction of stable phosphoranes Ph₃P:CHR with I. Moreover, an oxaphosphorin and oxazaphosphorin were prepared from the reaction of I with the phosphorane Ph₃P:CPh₂ and the iminophosphorane Ph₃P:NCO₂Et, resp.

IT 136829-50-0P 136848-91-4P
 RL: SPN (Synthetic preparation); PREP (Preparation)
 (preparation of)
 RN 136829-50-0 CAPLUS
 CN 1H-Indene-2-propanoic acid, β,β-dicyano-3-hydroxy-1-oxo-α-(triphenylphosphoranylidene)-, methyl ester (9CI) (CA INDEX NAME)



RN 136848-91-4 CAPLUS
 CN 1H-Indene-2-propanoic acid, β,β-dicyano-3-hydroxy-1-oxo-α-(triphenylphosphoranylidene)-, ethyl ester (9CI) (CA INDEX NAME)



L6 ANSWER 37 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1991:492003 CAPLUS

DOCUMENT NUMBER: 115:92003

TITLE: C-Alkylation of indoles with 1,1-bis(trifluoromethyl)-2,2-dicyanoethylene and 2-trifluoromethyl-3,3-dicyanoacrylic acid esters

AUTHOR(S): Chkanikov, N. D.; Komarov, K. V.; Tyutin, V. Yu.; Kolomiets, A. F.; Fokin, A. V.

CORPORATE SOURCE: Inst. Elementoorg. Soedin. im. Nesmeyanova, Moscow, USSR

SOURCE: Izvestiya Akademii Nauk SSSR, Seriya Khimicheskaya (1991), (5), 1193-5

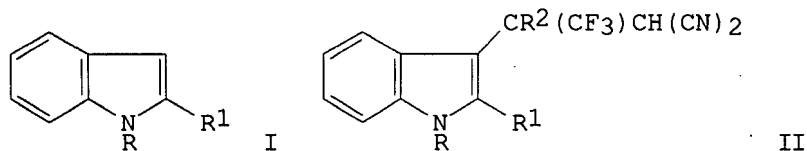
CODEN: IASKA6; ISSN: 0002-3353

DOCUMENT TYPE: Journal

LANGUAGE: Russian

OTHER SOURCE(S): CASREACT 115:92003

GI



AB The indole derivs. I (R = H, Me; R1 = H, Me, Ph) were alkylated with (NC)2C:CR2CF3 (R2 = CF3, CO2Me, CO2Et) to give the corresponding dicyanoethyl derivs. II.

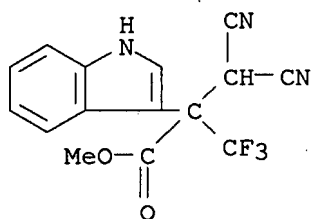
IT 135578-14-2P 135578-15-3P 135578-17-5P

135578-18-6P 135578-19-7P 135578-20-0P

RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of)

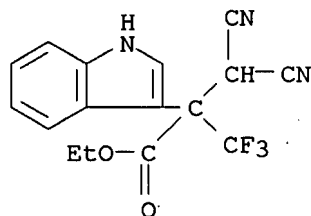
RN 135578-14-2 CAPLUS

CN 1H-Indole-3-acetic acid, α -(dicyanomethyl)- α -(trifluoromethyl)-, methyl ester (9CI) (CA INDEX NAME)



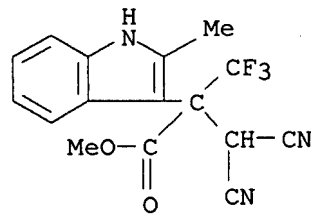
RN 135578-15-3 CAPLUS

CN 1H-Indole-3-acetic acid, α -(dicyanomethyl)- α -(trifluoromethyl)-, ethyl ester (9CI) (CA INDEX NAME)



RN 135578-17-5 CAPLUS

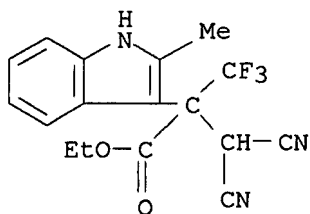
CN 1H-Indole-3-acetic acid, α -(dicyanomethyl)-2-methyl- α -(trifluoromethyl)-, methyl ester (9CI) (CA INDEX NAME)



RN 135578-18-6 CAPLUS

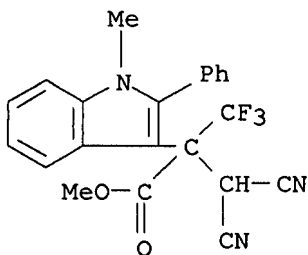
CN 1H-Indole-3-acetic acid, α -(dicyanomethyl)-2-methyl- α -

(trifluoromethyl)-, ethyl ester (9CI) (CA INDEX NAME)



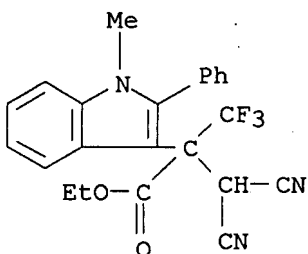
RN 135578-19-7 CAPLUS

CN 1H-Indole-3-acetic acid, α -(dicyanomethyl)-1-methyl-2-phenyl- α -(trifluoromethyl)-, methyl ester (9CI) (CA INDEX NAME)



RN 135578-20-0 CAPLUS

CN 1H-Indole-3-acetic acid, α -(dicyanomethyl)-1-methyl-2-phenyl- α -(trifluoromethyl)-, ethyl ester (9CI) (CA INDEX NAME)



L6 ANSWER 38 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1991:471633 CAPLUS

DOCUMENT NUMBER: 115:71633

TITLE: Preparation of pyrrolopyrimidines as antitumor agents

INVENTOR(S): Akimoto, Hiroshi; Hitaka, Takenori

PATENT ASSIGNEE(S): Takeda Chemical Industries, Ltd., Japan

SOURCE: Eur. Pat. Appl., 27 pp.

CODEN: EPXXDW

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

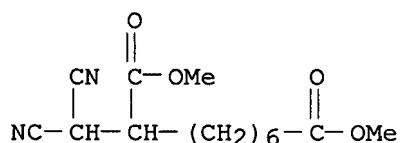
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 418924	A2	19910327	EP 1990-118202	19900921 <--

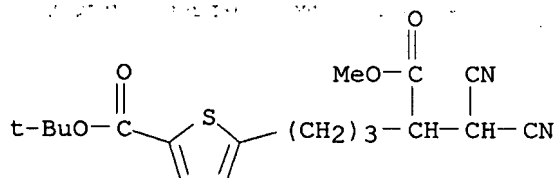
EP 418924 A3 19911023
 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE
 JP 03173890 A 19910729 JP 1990-249615 19900918 <--
 CA 2025830 A1 19910322 CA 1990-2025830 19900920 <--
 US 5354754 A 19941011 US 1993-46917 19930414 <--
 PRIORITY APPLN. INFO.: JP 1989-245998 A 19890921
 US 1990-585950 B1 19900921
 OTHER SOURCE(S): MARPAT 115:71633
 GI

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

AB Pyrrolopyrimidines [I; R1, R2 = H, ester residue; X = NH2, OH, SH; Y = H, OH; Z = (substituted) C2-4 divalent radical; Z1 = (substituted) divalent heterocycle residue, alkylene; dotted line indicates saturation or unsatn] are prepared Acetal II (1.32 g) (preparation given) was dissolved in CF3CO2H containing H2O with stirring at room temperature to give quant. salt III, which was dissolved with di-Et glutamate HCl in DMF and the solution was treated with 0.514 g H2NP(O)(OEt)2 and Et3N in DMF at room temperature to give 1.11 g diester IV (R1 = R2 = Et) (V). Saponification of 1.05 g V in THF gave 0.826 g acid IV (R1 = R2 = H), which showed IC50 of 0.00043 µg/mL against human epidermoid carcinoma KB cells.
 IT 135110-11-1P 135111-93-2P
 RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
 (preparation and reaction of, in preparation of antitumor agent)
 RN 135110-11-1 CAPLUS
 CN Nonanedioic acid, 2-(dicyanomethyl)-, dimethyl ester (9CI) (CA INDEX NAME)



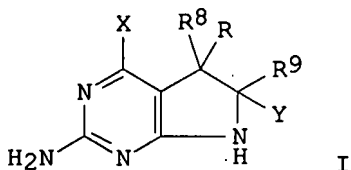
RN 135111-93-2 CAPLUS
 CN 2-Thiophenepentanoic acid, α-(dicyanomethyl)-5-[(1,1-dimethylethoxy)carbonyl]-, methyl ester (9CI) (CA INDEX NAME)



L6 ANSWER 39 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN
 ACCESSION NUMBER: 1991:429914 CAPLUS
 DOCUMENT NUMBER: 115:29914
 TITLE: Preparation of N-[[pyrrolopyrimidinylethyl]amino]benz

oyl]glutamates and analogs as antitumor agents
 INVENTOR(S): Akimoto, Hiroshi; Hitaka, Takenori; Miwa, Tetsuo
 PATENT ASSIGNEE(S): Takeda Chemical Industries, Ltd., Japan
 SOURCE: Eur. Pat. Appl., 51 pp.
 CODEN: EPXXDW
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 400562	A1	19901205	EP 1990-110131	19900529 <--
EP 400562	B1	19960821		
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE				
JP 04009382	A	19920114	JP 1990-136345	19900525 <--
JP 3015957	B2	20000306		
CA 2017604	A1	19901129	CA 1990-2017604	19900528 <--
AT 141603	T	19960915	AT 1990-110131	19900529 <--
PRIORITY APPLN. INFO.:			JP 1989-135642	A 19890529
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			JP 1990-93370	A 19900409
OTHER SOURCE(S):		MARPAT 115:29914		
GI				



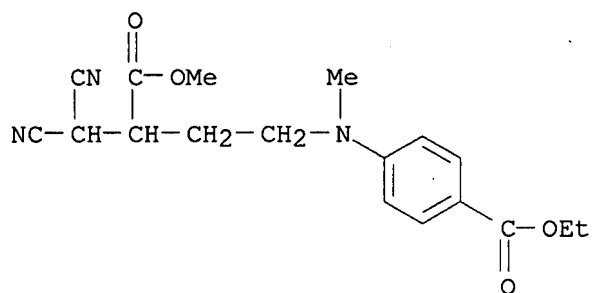
AB The title compds. [I; R = (CR1R2)iZ1(CR4R5)jZ2CONHCH(CO2R6)CH2CH2CO2R7; R1, R2, R4, R5 = H, hydrocarbyl, bond; R6, R7 = H, alkyl, (un)substituted Ph, PhCH2; R8, R9 = H; R8R9 = bond; X = NH2, OH, SH; Y = H, OH; Z1 = O, SOn, (alkyl)imino, etc.; Z2 = (un)substituted alkylene, divalent cyclic group; i, j = 0-3 (i + j = 1-3); n = 0-2] were prepared Thus, 4-(EtO2C)C6H4NMeCH2CH2CH(CO2Me)CH(CN)2 (preparation given) was cyclocondensed with guanidine and the product reduced to give, as 1 of 2 products, anilinoethylpyrrolopyrimidine I [R = CH2CH2NMeC6H4(COR10)-4; R8R9 = bond, X = NH2, Y = H] (II; R10 = OEt) which was condensed with di-Et L-glutamate to give, after saponification, L-II [R10 = NHCH(CO2H)CH2CH2CO2H]. The latter had

IC50 of 0.0013 μ M against human epidermoid carcinoma KB cell growth in vitro.

IT 133719-38-7P 133719-41-2P 133719-45-6P
 133719-47-8P
 RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
 (preparation and reaction of, in preparation of antitumor agents)

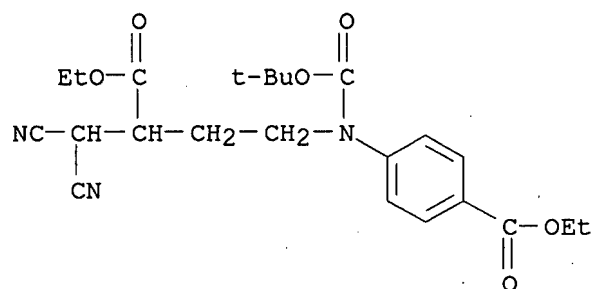
RN 133719-38-7 CAPLUS

CN Benzoic acid, 4-[[4,4-dicyano-3-(methoxycarbonyl)butyl]methyamino]-, ethyl ester (9CI) (CA INDEX NAME)



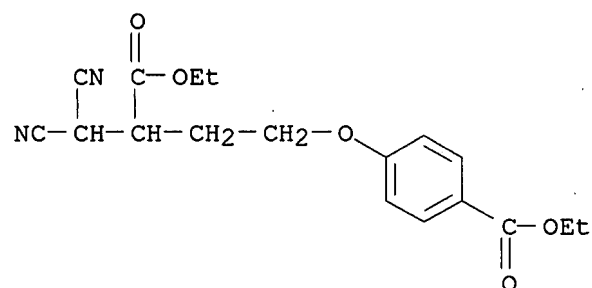
RN 133719-41-2 CAPLUS

CN Benzoic acid, 4-[[4,4-dicyano-3-(ethoxycarbonyl)butyl][(1,1-dimethylethoxy)carbonyl]amino]-, ethyl ester (9CI) (CA INDEX NAME)



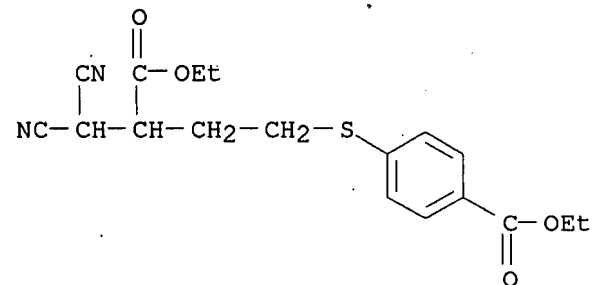
RN 133719-45-6 CAPLUS

CN Benzoic acid, 4-[4,4-dicyano-3-(ethoxycarbonyl)butoxy]-, ethyl ester (9CI) (CA INDEX NAME)



RN 133719-47-8 CAPLUS

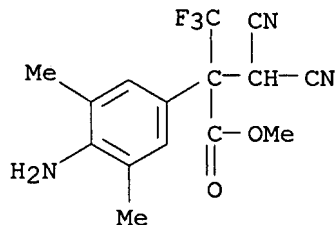
CN Benzoic acid, 4-[[4,4-dicyano-3-(ethoxycarbonyl)butyl]thio]-, ethyl ester (9CI) (CA INDEX NAME)



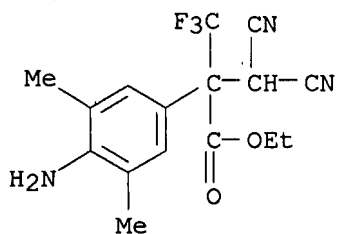
L6 ANSWER 40 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN
 ACCESSION NUMBER: 1991:429194 CAPLUS
 DOCUMENT NUMBER: 115:29194
 TITLE: Synthesis of esters of 3,3-dicyano-2-(trifluoromethyl)acrylic acid and their reactions with aryl amines
 AUTHOR(S): Tyutin, V. Y.; Chkanikov, N. D.; Kolomiets, A. F.; Fokin, A. V.
 CORPORATE SOURCE: Inst. Organoelem. Compd., Moscow, 117813, USSR
 SOURCE: Journal of Fluorine Chemistry (1991), 51(3), 323-34
 CODEN: JFLCAR; ISSN: 0022-1139
 DOCUMENT TYPE: Journal
 LANGUAGE: English
 OTHER SOURCE(S): CASREACT 115:29194
 GI

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

AB Title acrylates (NC)₂C:C(CF₃)CO₂R (I; R = Me, Et) were prepared by the condensation of CH₂(CN)₂ with CF₃COCO₂R in presence of ZnCl₂. Reaction of I with aromatic amines was investigated. Thus, 2,6-dimethylaniline reacted with I in CHCl₃ to give adduct II. 2,5-Dimethoxyaniline, and Ph₂NH gave similar adducts. o- And m-C₆H₄(NH₂)₂ reacted with I to give cyclocondensation products, quinoxalinone III and indoline IV resp. 4-R₁C₆H₄NH₂ (R₁ = H, NO₂) gave pyrazolines V on cyclocondensation with I. Reaction of I with 3-aminopyrazole gave pyrazolopyridines VI.
 IT 134641-38-6P 134641-39-7P 134641-40-0P
 134641-41-1P 134641-42-2P 134641-43-3P
 134641-44-4P
 RL: SPN (Synthetic preparation); PREP (Preparation) (preparation of)
 RN 134641-38-6 CAPLUS
 CN Benzeneacetic acid, 4-amino-α-(dicyanomethyl)-3,5-dimethyl-α-(trifluoromethyl)-, methyl ester (9CI) (CA INDEX NAME)

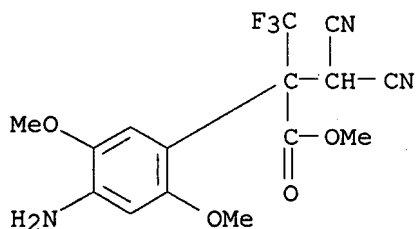


RN 134641-39-7 CAPLUS
 CN Benzeneacetic acid, 4-amino-α-(dicyanomethyl)-3,5-dimethyl-α-(trifluoromethyl)-, ethyl ester (9CI) (CA INDEX NAME)



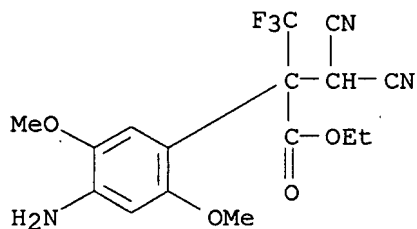
RN 134641-40-0 CAPLUS

CN Benzeneacetic acid, 4-amino-α-(dicyanomethyl)-2,5-dimethoxy-α-(trifluoromethyl)-, methyl ester (9CI) (CA INDEX NAME)



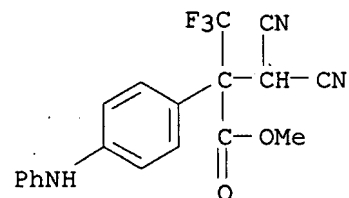
RN 134641-41-1 CAPLUS

CN Benzeneacetic acid, 4-amino-α-(dicyanomethyl)-2,5-dimethoxy-α-(trifluoromethyl)-, ethyl ester (9CI) (CA INDEX NAME)



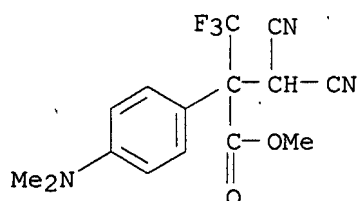
RN 134641-42-2 CAPLUS

CN Benzeneacetic acid, α-(dicyanomethyl)-4-(phenylamino)-α-(trifluoromethyl)-, methyl ester (9CI) (CA INDEX NAME)

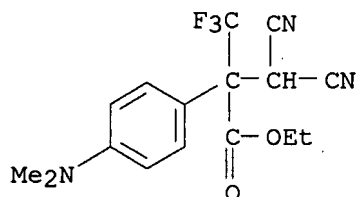


RN 134641-43-3 CAPLUS

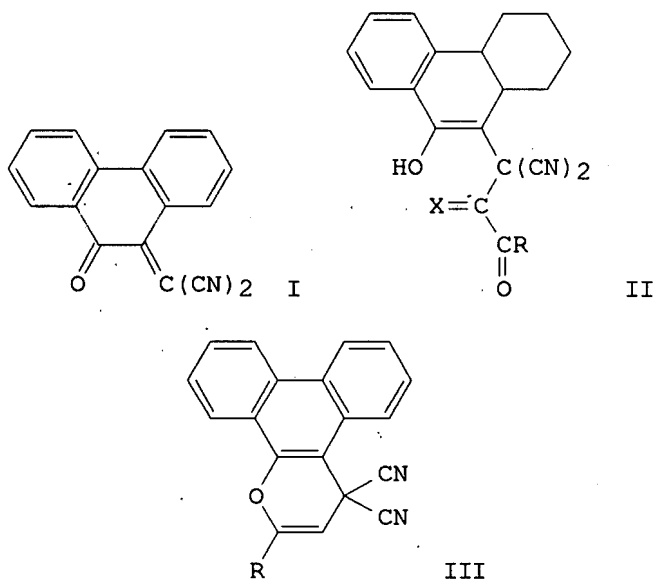
CN Benzeneacetic acid, α-(dicyanomethyl)-4-(dimethylamino)-α-(trifluoromethyl)-, methyl ester (9CI) (CA INDEX NAME)



RN 134641-44-4 CAPLUS
 CN Benzeneacetic acid, α -(dicyanomethyl)-4-(dimethylamino)- α -(trifluoromethyl)-, ethyl ester (9CI) (CA INDEX NAME)



L6 ANSWER 41 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN
 ACCESSION NUMBER: 1991:247080 CAPLUS
 DOCUMENT NUMBER: 114:247080
 TITLE: Reaction of α,β -unsaturated nitriles with phosphorus ylides
 AUTHOR(S): Abdou, Wafaa M.; Ganoub, Neven A. F.
 CORPORATE SOURCE: Natl. Res. Cent., Dokki, Egypt
 SOURCE: Chemistry & Industry (London, United Kingdom) (1991), (6), 217-18
 CODEN: CHINAG; ISSN: 0009-3068
 DOCUMENT TYPE: Journal
 LANGUAGE: English
 OTHER SOURCE(S): CASREACT 114:247080
 GI

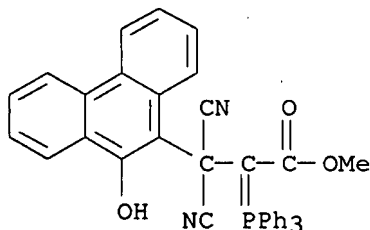


AB Reaction of unsatd. nitrile I with ROCC-HP+Ph₃ (R = OMe, OEt, Ph) in benzene at 25° gave 75-80% addition products II (X = PPh₃). On heating II (X = PPh₃) to 200° they underwent an intramol. Wittig reaction to give arenopyrans III. Heating phosphorane II (R = OMe, X = PPh₃) with BzH gave II (X = CHPh).

IT 133973-19-0P 133973-20-3P
 RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
 (preparation and intramol. Wittig reaction of)

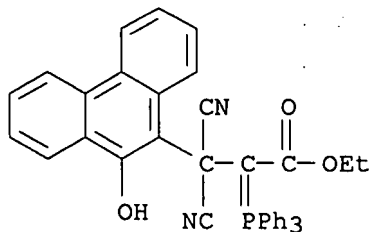
RN 133973-19-0 CAPLUS

CN 9-Phenanthrenepropanoic acid, β,β-dicyano-10-hydroxy-α-(triphenylphosphoranylidene)-, methyl ester (9CI) (CA INDEX NAME)



RN 133973-20-3 CAPLUS

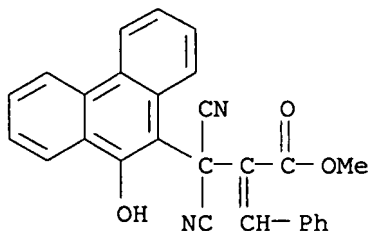
CN 9-Phenanthrenepropanoic acid, β,β-dicyano-10-hydroxy-α-(triphenylphosphoranylidene)-, ethyl ester (9CI) (CA INDEX NAME)



IT 133973-25-8P
 RL: SPN (Synthetic preparation); PREP (Preparation)
 (preparation of)

RN 133973-25-8 CAPLUS

CN 9-Phenanthrenepropanoic acid, β,β-dicyano-10-hydroxy-α-(phenylmethylene)-, methyl ester (9CI) (CA INDEX NAME)



TITLE: Reactions of malononitrile with acetylenic esters and ketones [Erratum to document cited in CA113(25):231170y]

AUTHOR(S): Kandeel, Kamal A.; Vernon, John M.; Dransfield, Trevor A.; Fouli, Fouli A.; Youssef, Ahmed S. A.

CORPORATE SOURCE: Dep. Chem., Univ. York, Heslington/York, YO1 5DD, UK

SOURCE: Journal of Chemical Research, Synopses (1990), (12), 406
CODEN: JRPSDC; ISSN: 0308-2342

DOCUMENT TYPE: Journal

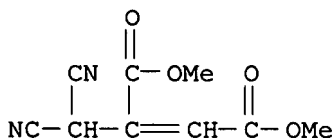
LANGUAGE: English

AB An error in the structure for compound 13 has been corrected The error was not reflected in the abstract or the index entries.

IT 130747-61-4P
RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of (Erratum))

RN 130747-61-4 CAPLUS

CN 2-Butenedioic acid, 2-(dicyanomethyl)-, dimethyl ester (9CI) (CA INDEX NAME)



L6 ANSWER 43 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1991:82446 CAPLUS

DOCUMENT NUMBER: 114:82446

TITLE: Novel pyrrolo[2,3-d]pyrimidine antifolates: synthesis and antitumor activities

AUTHOR(S): Miwa, Tetsuo; Hitaka, Takenori; Akimoto, Hiroshi; Nomura, Hiroaki

CORPORATE SOURCE: Res. Dev. Div., Takeda Chem. Ind., Ltd., Osaka, 532, Japan

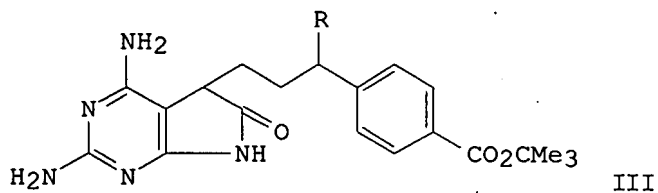
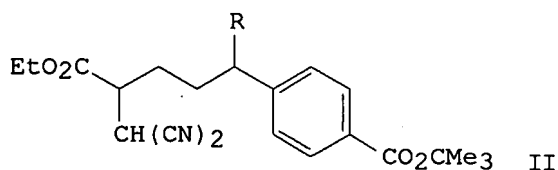
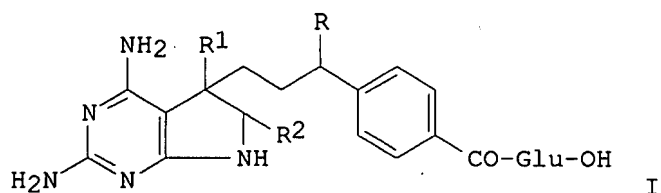
SOURCE: Journal of Medicinal Chemistry (1991), 34(2), 555-60
CODEN: JMCMAR; ISSN: 0022-2623

DOCUMENT TYPE: Journal

LANGUAGE: English

OTHER SOURCE(S): CASREACT 114:82446

GI



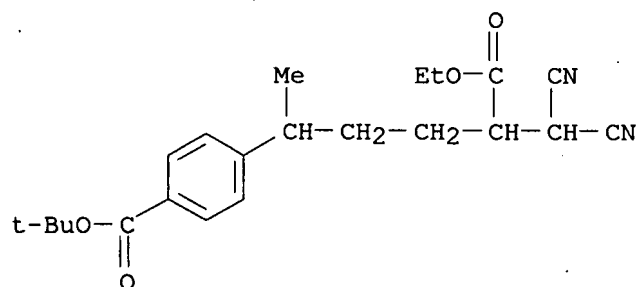
AB Title compds. I (R = H, Me; R1 = H, Et; R2R3 = bond, R2 = R3 = H) were prepared as antifolates. A key step was the cyclocondensation of dicyano compound II (R = H, Me) with guanidine-HCl to give pyrrolo[2,3-d]pyrimidines III. III were prepared in several steps from p-RCOC6H4CO2CMe2 and CH3CH:CHCO2Et or BrCH2CH:CHCO2Et. These antifolates were more growth-inhibitory by about 1 order of magnitude than methotrexate (MTX) against KB human epidermoid carcinoma cells and A549 human nonsmall cell lung carcinoma cells in in vitro culture.

IT 125991-47-1P 130351-33-6P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation and cyclocondensation of, with guanidine)

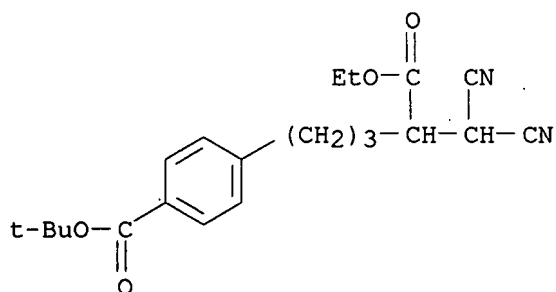
RN 125991-47-1 CAPLUS

CN Benzenepentanoic acid, α -(dicyanomethyl)-4-[(1,1-dimethylethoxy)carbonyl]- δ -methyl-, ethyl ester (9CI) (CA INDEX NAME)

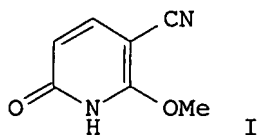


RN 130351-33-6 CAPLUS

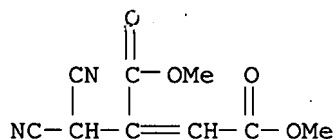
CN Benzenepentanoic acid, α -(dicyanomethyl)-4-[(1,1-dimethylethoxy)carbonyl]-, ethyl ester (9CI) (CA INDEX NAME)



L6 ANSWER 44 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN
 ACCESSION NUMBER: 1990:631170 CAPLUS
 DOCUMENT NUMBER: 113:231170
 TITLE: Reactions of malononitrile with acetylenic esters and ketones
 AUTHOR(S): Kandeel, Kamal A.; Vernon, John M.; Dransfield, Trevor A.; Fouli, Fouli A.; Youssef, Ahmed S. A.
 CORPORATE SOURCE: Dep. Chem., Univ. York, Heslington/York, YO1 5DD, UK
 SOURCE: Journal of Chemical Research, Synopses (1990), (9), 276-7
 CODEN: JRPSDC; ISSN: 0308-2342
 DOCUMENT TYPE: Journal
 LANGUAGE: English
 OTHER SOURCE(S): CASREACT 113:231170
 GI

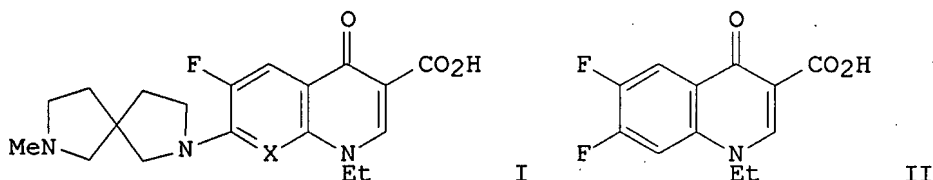


AB The addition of malononitrile to acetylenic esters and acetylenic ketones catalyzed by sodium alkoxides gave 3- and 5-cyano-2-pyridones, e.g., I, 2-cyano- and 2,6-dicyanoaniline, and other products.
 IT 130747-61-4P
 RL: SPN (Synthetic preparation); PREP (Preparation) (preparation of)
 RN 130747-61-4 CAPLUS
 CN 2-Butenedioic acid, 2-(dicyanomethyl)-, dimethyl ester (9CI) (CA INDEX NAME)

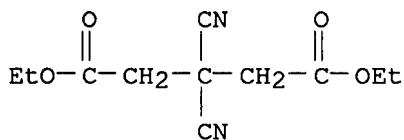


L6 ANSWER 45 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN
 ACCESSION NUMBER: 1990:497432 CAPLUS
 DOCUMENT NUMBER: 113:97432
 TITLE: Quinolone antibacterial agents substituted at the 7-position with spiroamines. Synthesis and

structure-activity relationships
 AUTHOR(S): Culbertson, Townley P.; Sanchez, Joseph P.; Gambino, Laura; Sesnie, Josephine A.
 CORPORATE SOURCE: Parke-Davis Pharm. Res. Div., Warner-Lambert Co., Ann Arbor, MI, 48105, USA
 SOURCE: Journal of Medicinal Chemistry (1990), 33(8), 2270-5
 CODEN: JMCMAR; ISSN: 0022-2623
 DOCUMENT TYPE: Journal
 LANGUAGE: English
 OTHER SOURCE(S): CASREACT 113:97432
 GI



AB Fluoroquinolone antibacterials having the 7-position (10-position of pyridobenzoxazines) substituted with 2,7-diazaspiro[4.4]nonane, 1,7-diazaspiro[4.4]nonane, or 2,8-diazaspiro[5.5]undecane (e.g. I. (X = CF, CH, N) were prepared and their biol. activities were compared with piperazine and pyrrolidine substituted analogs. Most exhibited potent Gram-pos. and Gram-neg. activity, especially when side chain was N-alkylated. Thus, the quinolinecarboxylic acid II was treated with 2-methyl-2,7-diazaspiro[4.4]nonane to give I (X = CH).
 IT 77415-69-1
 RL: RCT (Reactant); RACT (Reactant or reagent)
 (reductive cyclization of)
 RN 77415-69-1 CAPLUS
 CN Pentanedioic acid, 3,3-dicyano-, diethyl ester (9CI) (CA INDEX NAME)



L6 ANSWER 46 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN
 ACCESSION NUMBER: 1990:235871 CAPLUS
 DOCUMENT NUMBER: 112:235871
 TITLE: New gem-dicyanocyclobutane-containing hydroxyesters
 AUTHOR(S): Mori, Shoji; Kakuchi, Toyoji; Padias, Anne Buyle; Hall, H. K., Jr.
 CORPORATE SOURCE: Chem. Dep., Univ. Arizona, Tucson, AZ, 85721, USA
 SOURCE: Journal of Polymer Science, Part A: Polymer Chemistry (1990), 28(3), 551-8
 CODEN: JPACEC; ISSN: 0887-624X
 DOCUMENT TYPE: Journal
 LANGUAGE: English
 AB Six gem-dicyanocyclobutanes containing carbomethoxy and hydroxyl/acetoxyl functions were synthesized by cycloaddn. of the appropriate vinyl ethers

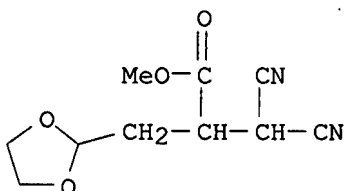
or alkoxytyrenes to Me β,β -dicyanoacrylate. They were too thermally liable to allow polycondensation to potentially piezoelec. linear polyesters.

IT 127396-28-5P

RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation and attempted polymerization of)

RN 127396-28-5 CAPLUS

CN 1,3-Dioxolane-2-propanoic acid, α -(dicyanomethyl)-, methyl ester
(9CI) (CA INDEX NAME)



L6 ANSWER 47 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1990:158968 CAPLUS

DOCUMENT NUMBER: 112:158968

TITLE: Preparation of N-[(pyrrolopyrimidinylalkyl)benzoyl]glutamates as neoplasm inhibitors

INVENTOR(S): Akimoto, Hiroshi; Hitaka, Takenori; Miwa, Tetsuo

PATENT ASSIGNEE(S): Takeda Chemical Industries, Ltd., Japan

SOURCE: Eur. Pat. Appl., 24 pp.

CODEN: EPXXDW

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

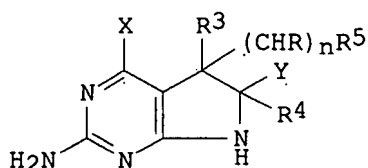
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 334636	A2	19890927	EP 1989-302851	19890322 <--
EP 334636	A3	19910502		
EP 334636	B1	19961023		
R: AT, BE, CH, DE, ES, FR, GB, GR, IT, LI, LU, NL, SE				
NO 8901206	A	19890925	NO 1989-1206	19890320 <--
NO 169490	B	19920323		
NO 169490	C	19920701		
US 4997838	A	19910305	US 1989-326901	19890321 <--
DK 8901437	A	19890925	DK 1989-1437	19890322 <--
DK 173980	B1	20020325		
AT 144513	T	19961115	AT 1989-302851	19890322 <--
ES 2092994	T3	19961216	ES 1989-302851	19890322 <--
CA 1340794	C	19991019	CA 1989-594699	19890323 <--
CN 1037513	A	19891129	CN 1989-101681	19890324 <--
CN 1029970	B	19951011		
HU 51624	A2	19900528	HU 1989-1517	19890324 <--
HU 203105	B	19910528		
JP 02167281	A	19900627	JP 1989-72235	19890324 <--
JP 07005599	B	19950125		
HU 55396	A2	19910528	HU 1990-8458	19890324 <--
HU 215928	B	19990329		
US 5106974	A	19920421	US 1990-578258	19900906 <--
NO 9100661	A	19890925	NO 1991-661	19910219 <--
NO 178304	B	19951120		
NO 178304	C	19960228		

US 5296600
US 5539113
PRIORITY APPLN. INFO.:

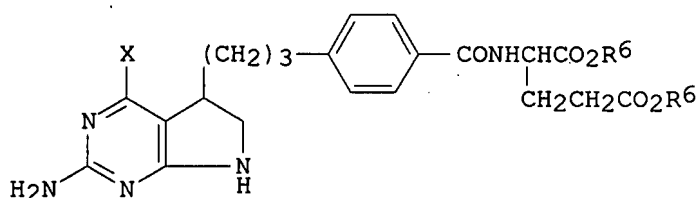
A 19940322
A 19960723

US 1992-824106 19920122 <--
US 1993-161533 19931206 <--
JP 1988-71149 A 19880324
JP 1988-245379 A 19880929
NO 1989-1206 A1 19890320
US 1989-326901 A3 19890321
US 1990-578258 A3 19900906
US 1992-824106 A3 19920122

OTHER SOURCE(S): CASREACT 112:158968; MARPAT 112:158968
GI

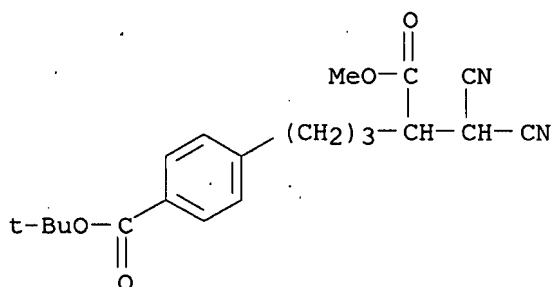


I



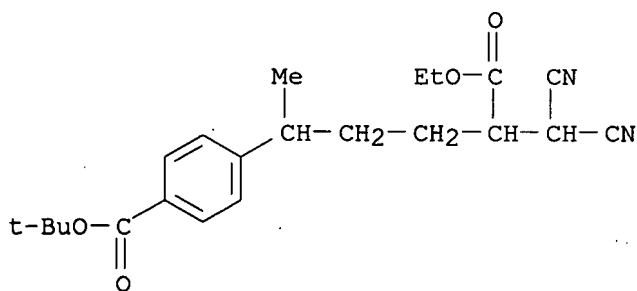
III

- AB The title compds. [I; R = H, F, alkyl, alkenyl, alkynyl; R3, R4 = H; R3R4 = bond; R5 = C6H4(CONHCHR1CH2CH2R2)-4; R1,R2 = (un)esterified CO2H; X = NH2, OH; Y = H, NH2, OH; n = 2-4] were prepared Thus, 4-(Me3CO2C)C6H4(CH2)3CH[CH(CN)2]CO2Me (preparation given) was refluxed 28 h with (H2N)2C:NH.HCl in Me3COH containing Me3COK to give I [R = R3 = H, R5 = C6H4(CO2CMe3)-4, X = NH2, n = 3] (II; R4Y = O) which was hydrogenated to II (R4 = Y = H). The latter was hydrolyzed and the product condensed with di-Et L-glutamate to give title compound III (R6 = Et, X = NH2) which was hydrolyzed to III (R6 = H, X = OH) which had IC50 of 0.0006 µg/mL against human nasopharyngeal cancer KB cells in vitro.
- IT 125991-38-0P 125991-47-1P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation and reaction of, in preparation of neoplasm inhibitors)
- RN 125991-38-0 CAPLUS
- CN Benzenepentanoic acid, α-(dicyanomethyl)-4-[(1,1-dimethylethoxy)carbonyl]-, methyl ester (9CI) (CA INDEX NAME)



- RN 125991-47-1 CAPLUS
- CN Benzenepentanoic acid, α-(dicyanomethyl)-4-[(1,1-

dimethylethoxy)carbonyl]-8-methyl-, ethyl ester (9CI) (CA INDEX NAME)



L6 ANSWER 48 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN
 ACCESSION NUMBER: 1989:415389 CAPLUS
 DOCUMENT NUMBER: 111:15389
 TITLE: Color photothermographic elements containing leuco compounds
 INVENTOR(S): Sakizadeh, Kumars; Weigel, David C.; Grieve, Duncan; Poon, Stephen S. C.; Thien, Tran V.
 PATENT ASSIGNEE(S): Minnesota Mining and Manufacturing Co., USA
 SOURCE: Eur. Pat. Appl., 35 pp.
 CODEN: EPXXDW
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 294099	A2	19881207	EP 1988-304771	19880526 <--
EP 294099	A3	19890531		
EP 294099	B1	19930818		
R: BE, DE, FR, GB, IT				
US 4883747	A	19891128	US 1988-200665	19880531 <--
CA 1331107	C	19940802	CA 1988-568396	19880602 <--
AU 8817345	A	19881208	AU 1988-17345	19880603 <--
AU 606162	B2	19910131		
JP 02032332	A	19900202	JP 1988-145566	19880613 <--
JP 2590204	B2	19970312		
US 4923792	A	19900508	US 1989-368566	19890620 <--
PRIORITY APPLN. INFO.:			GB 1987-12961	A 19870603
			US 1988-200665	A1 19880531

OTHER SOURCE(S): CASREACT 111:15389; MARPAT 111:15389

AB A photothermog. material comprises Ag halide in reactive association with a Ag salt of an organic acid and a color-generating reducing agent which is a leuco compound oxidizable by Ag ions into a colored dye of the formula $ArR_1C(:C(R_5)C(R_4):)_nCR_2R_3$ [$n = 0-2$; $R_1 = H, CN, C1-5$ alkyl, aryl, CO_2R_6 ; $R_6 = C1-5$ alkyl or aryl; $R_2, R_3 = CN, NO_2, CO_2R_6, SO_2R_6, COR_6$; R_3 and R_2 may combine together to form a ring; $R_4, R_5 = H, CN, C1-5$ alkyl, or R_4 and R_5 together may form a ring; $Ar =$ thienyl, furyl, phenyl]. The material produces images with improved color stability. Thus, a green-yellow image was produced with a photothermog. material incorporating leuco form of (p-dimethylaminobenzylidene)dimethylbarbituric acid.

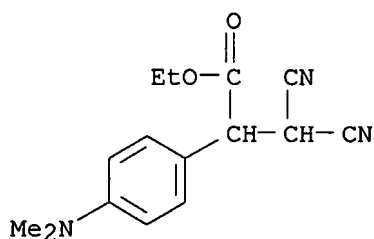
IT 121246-61-5

RL: USES (Uses)

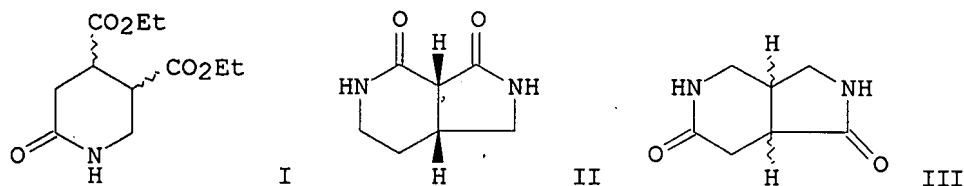
(photothermog. material containing, for improved image stability)

RN 121246-61-5 CAPLUS

CN Benzeneacetic acid, α -(dicyanomethyl)-4-(dimethylamino)-, ethyl ester (9CI) (CA INDEX NAME)



L6 ANSWER 49 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN
 ACCESSION NUMBER: 1988:570272 CAPLUS
 DOCUMENT NUMBER: 109:170272
 TITLE: Synthesis and cognition-activating properties of some mono- and bicyclic lactam derivatives
 AUTHOR(S): Altomare, Cosimo; Carotti, Angelo; Casini, Giovanni; Cellamare, Saverio; Ferappi, Marcello; Gavuzzo, Enrico; Mazza, Fernando; Pantaleoni, Giancarlo; Giorgi, Raffaele
 CORPORATE SOURCE: Dip. Farm.-Chim., Univ. Bari, Bari, Italy
 SOURCE: Journal of Medicinal Chemistry (1988), 31(11), 2153-8
 CODEN: JMCMAR; ISSN: 0022-2623
 DOCUMENT TYPE: Journal
 LANGUAGE: English
 OTHER SOURCE(S): CASREACT 109:170272
 GI



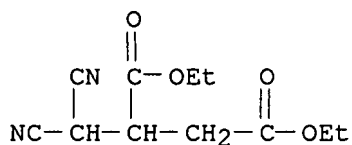
AB Upon reductive cyclization cyano esters $\text{EtO}_2\text{CCH}_2\text{CH}(\text{CO}_2\text{Et})\text{CHRCN}$ ($\text{R} = \text{CO}_2\text{Et}$, cyano) and $\text{NCCH}_2\text{CH}(\text{CN})\text{CH}(\text{CO}_2\text{Et})_2$ yielded piperidones and perhydropyrrolo[3,4-c]pyridine lactams I, II and III, resp. generally as a mixture of diastereomeric cis-trans forms. X-ray crystallog. anal. were carried out on cis-II and III. A series of neuropsychopharmacol. tests performed on I, II; and III indicated that they are generally nontoxic even at high doses (up to 1000 mg/kg i.p.)⁹. The cognition activating properties of lactams cis- and trans-I, cis-II, and III were evaluated in enhancing retention for passive avoidance learning in rats without and after electroconvulsive shock (ECS); compds. cis-I and III were found to be more potent than piracetam in the amnesia-reversal testing.

IT 82584-86-9

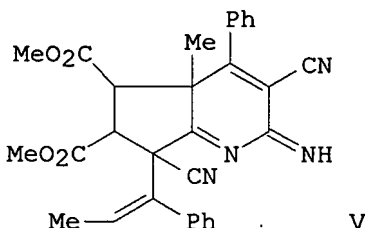
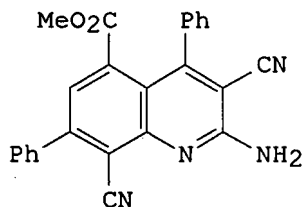
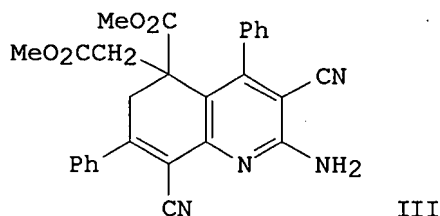
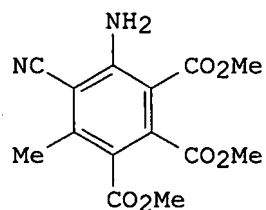
RL: RCT (Reactant); RACT (Reactant or reagent)
 (reductive cyclization of, cyclic lactams from)

RN 82584-86-9 CAPLUS

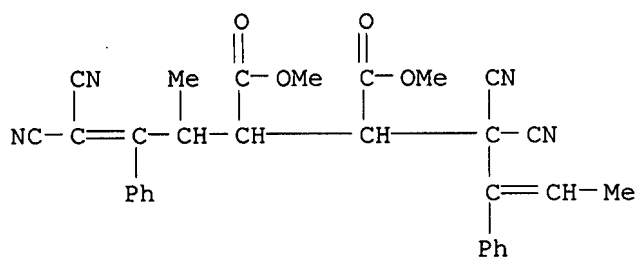
CN Butanedioic acid, (dicyanomethyl)-, diethyl ester (9CI) (CA INDEX NAME)



L6 ANSWER 50 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN
 ACCESSION NUMBER: 1988:94172 CAPLUS
 DOCUMENT NUMBER: 108:94172
 TITLE: Addition of ylidenemalononitriles onto dimethyl
 acetylenedicarboxylate
 AUTHOR(S): Gewalt, Karl; Hain, Ute; Gruner, Margit
 CORPORATE SOURCE: Sekt. Chem., Tech. Univ. Dresden, Dresden, DDR-8027,
 Ger. Dem. Rep.
 SOURCE: Zeitschrift fuer Chemie (1987), 27(1), 32-4
 CODEN: ZECEAL; ISSN: 0044-2402
 DOCUMENT TYPE: Journal
 LANGUAGE: German
 OTHER SOURCE(S): CASREACT 108:94172
 GI



AB (NC)2C:CMech2CO2Me and MeO2CC.tplbond.CCO2Me (I) in the presence of K2CO3
 cycloadded to give 49% aniline II. (NC)2C:CPhMe and I, treated with Et3N,
 gave 30% dihydroquinoline III, which was aromatized by heating at
 270° in N₂-MeOH to 80% quinolinecarboxylate IV. (NC)2C:CPhEt and I
 gave 29% (NC)2C:CPhCHMeCH(CO2Me)CH(CO2Me)C(CN)2CPh:CHMe, which cyclized to
 cyclopentapyridinedicarboxylate V.
 IT 112754-03-7P
 RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
 (Reactant or reagent)
 (preparation and thermal intramol. cyclization of)
 RN 112754-03-7 CAPLUS
 CN Butanedioic acid, 2-(3,3-dicyano-1-methyl-2-phenyl-2-propenyl)-3-(1,1-
 dicyano-2-phenyl-2-butenyl)-, dimethyl ester (9CI) (CA INDEX NAME)



L6 ANSWER 51 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1987:597249 CAPLUS

DOCUMENT NUMBER: 107:197249

TITLE: Influence of the solvent on the nature of a tetramethylene biradical intermediate.

AUTHOR(S): Padias, Anne Buyle; Hall, H. K., Jr.

CORPORATE SOURCE: Chem. Dep., Univ. Arizona, Tucson, AZ, 85721, USA

SOURCE: Journal of Organic Chemistry (1987), 52(20), 4536-9

CODEN: JOCEAH; ISSN: 0022-3263

DOCUMENT TYPE: Journal

LANGUAGE: English

OTHER SOURCE(S): CASREACT 107:197249

AB In the spontaneous thermal reactions of p-methoxystyrene and Me 3,3-dicyanoacrylate, several reaction products are observed: a 1/1 alternating copolymer, a double Diels-Alder adduct, and the cyclobutane adduct. In dipolar aprotic solvents, no polymerization occurs, and the double Diels-Alder adduct is favored; in protic polar solvents cyclobutane formation competes with copolymer. In nonpolar solvents, copolymer dominates. A biradical tetramethylene species is proposed as the key intermediate. In polar solvents, this biradical exhibits considerable polar character, and Coulombic attraction between the termini favors the coiled or gauche conformation, leading preferentially to cycloadducts. In nonpolar solvents, the trans conformation initiates the polymerization. The

main factors influencing the products are the solvent polarity and the ability of the solvent to interact with the biradical.

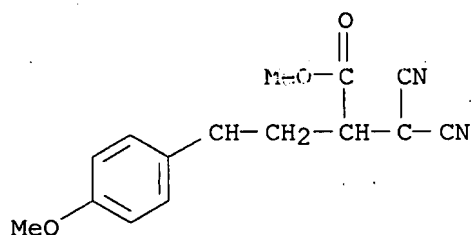
IT 110193-00-5

RL: PRP (Properties)

(conformation and spin and electron d. of, solvent effects on)

RN 110193-00-5 CAPLUS

CN 1,4-Butanediyl, 1,1-dicyano-2-(methoxycarbonyl)-4-(4-methoxyphenyl)- (9CI)
(CA INDEX NAME)



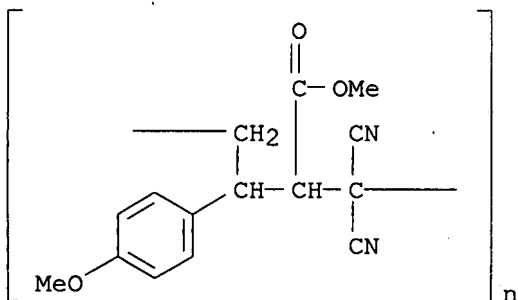
IT 110193-05-0P

RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of)

RN 110193-05-0 CAPLUS

CN Poly[1,1-dicyano-2-(methoxycarbonyl)-3-(4-methoxyphenyl)-1,4-butanediyl]

(9CI) (CA INDEX NAME)



L6 ANSWER 52 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1987:196834 CAPLUS

DOCUMENT NUMBER: 106:196834

TITLE: Cationic polymerization of nitrogen-containing electron-rich vinyl monomers by electrophilic olefins and their cyclobutane cycloadducts

AUTHOR(S): Abdelkader, Mohamed; Padias, Anne Buyle; Hall, H. K., Jr.

CORPORATE SOURCE: Chem. Dep., Univ. Arizona, Tucson, AZ, 85721, USA

SOURCE: Macromolecules (1987), 20(5), 944-8

CODEN: MAMOBX; ISSN: 0024-9297

DOCUMENT TYPE: Journal

LANGUAGE: English

AB The major pathways for the reactions of very electron-rich N-containing olefins with several electrophilic olefins were studied. N-Ethyl-3-vinylcarbazole (I) [1486-07-3], N-vinylcarbazole (II) [1484-13-5], and p-(dimethylamino)styrene (III) [2039-80-7] underwent kinetic cyclobutane formation with an electrophilic olefin without a leaving group, Me β , β -dicyanoacrylate (IV) [82849-50-1], and one with a weak β -leaving group, tetracyanoethylene (V) [670-54-2]. The third electrophilic olefin, β , β -dicyanovinyl chloride (VI) [10472-09-0], had a strong β -leaving group and readily initiated the cationic polymerization of I and II and oligomerization of III. If an excess

of

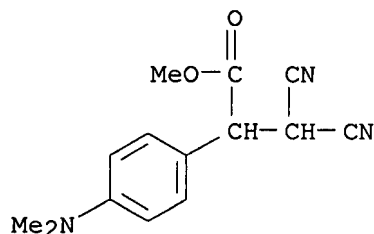
donor olefin was used, IV, V, and VI all initiated cationic homopolymerization of I and II, while III only led to oligomers, as it did with conventional Brønsted initiators. Cationic initiation by their own cyclobutane adducts was observed for the very electron-rich monomers I and II. Postcyanovinylation of the formed polymers by the electrophilic olefins occurred. Incorporation of a β -leaving group enhanced the initiating ability of the electrophilic olefins and N-carbazyl and N-ethyl-3-carbazyl were overall the most effective donor substituents favoring cationic homopolymerization.

IT 107540-79-4F

RL: SPN (Synthetic preparation); PREP. (Preparation)
(preparation of, from dimethylaniline and dicyanovinyl compound)

RN 107540-79-4 CAPLUS

CN Benzeneacetic acid, α -(dicyanomethyl)-4-(dimethylamino)-, methyl ester (9CI) (CA INDEX NAME)



L6 ANSWER 53 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN
 ACCESSION NUMBER: 1987:33439 CAPLUS
 DOCUMENT NUMBER: 106:33439
 TITLE: Synthesis of novel symmetric diamino acids
 AUTHOR(S): Reddy, P. Anantha; Erickson, Bruce W.
 CORPORATE SOURCE: Rockefeller Univ., New York, NY, 10021, USA
 SOURCE: Pept.: Struct. Funct., Proc. Am. Pept. Symp., 9th (1985), 453-6
 CODEN: 54ZNAJ

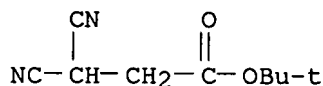
DOCUMENT TYPE: Conference
 LANGUAGE: English

AB Sym diamino acids (H₂NCH₂)₂CHCH₂CO₂H (Aab) 3,5-(H₂NCH₂X)₂C₆H₃CO₂H [X = null, CH₂ (Bab)] were prepared from (NC)₂CHCH₂CO₂CMe₃ and 3,5-(BrCH₂)₂C₆H₃CO₂Me. The N,N-bis(tert-butoxycarbonyl) derivative of Aab couples efficiently during solid-phase peptide synthesis. The corresponding derivative of Bab is used in the synthesis of the protein betabellin.

IT 105995-37-7P 105995-39-9P
 RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
 (preparation and hydrogenation of)

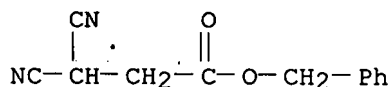
RN 105995-37-7 CAPLUS

CN Propanoic acid, 3,3-dicyano-, 1,1-dimethylethyl ester (9CI) (CA INDEX NAME)



RN 105995-39-9 CAPLUS

CN Propanoic acid, 3,3-dicyano-, phenylmethyl ester (9CI) (CA INDEX NAME)



L6 ANSWER 54 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1986:496793 CAPLUS

DOCUMENT NUMBER: 105:96793

TITLE: Zwitterionic tetramethylenes as the common intermediates in the cycloaddition and polymerization reactions of N-vinylcarbazole with electrophilic tetrasubstituted ethylenes: a new explanation for charge-transfer initiation

AUTHOR(S): Gotoh, Tetsuya; Padias, Anne Buyle; Hall, H. K., Jr.

CORPORATE SOURCE: Chem. Dep., Univ. Arizona, Tucson, AZ, 85721, USA
 SOURCE: Journal of the American Chemical Society (1986
), 108(16), 4920-31
 CODEN: JACSAT; ISSN: 0002-7863
 DOCUMENT TYPE: Journal
 LANGUAGE: English
 OTHER SOURCE(S): CASREACT 105:96793

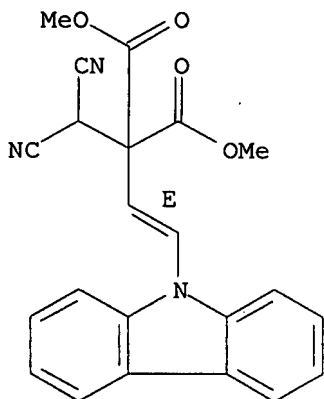
AB The reactions of N-vinylcarbazole (I) with electrophilic tetrasubstituted ethylenes were examples of reactions whose outcomes are manipulated by changes in concentration, structure, and working procedure to form either small mols. (cyclobutanes, 1-butenes) or poly(vinylcarbazole). Equivalent concns. and evaporative workup (organic chemists' conditions) lead to small mols.; a large excess of I and precipitative workup give polymer. The mechanism involves gauche and trans zwitterionic tetramethylenes as intermediates. The former gives cyclobutane reversibly. The latter gives 1-butenes intramol. or adds monomers to form cyclohexanes or eventually polymer. The organic chemical and polymer chemical are unified on this basis. Extensive stereochem. and kinetic support for these propositions is given. Two other proposed mechanisms for these charge-transfer initiations are excluded.

IT 96735-90-9P 102852-12-0P 102852-13-1P
 102852-14-2P 102852-38-0P 102852-39-1P
 RL: SPN (Synthetic preparation); PREP (Preparation)
 (preparation of)

RN 96735-90-9 CAPLUS

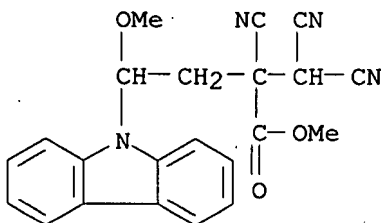
CN Propanedioic acid, [2-(9H-carbazol-9-yl)ethenyl] (dicyanomethyl)-, dimethyl ester, (E)- (9CI) (CA INDEX NAME)

Double bond geometry as shown.



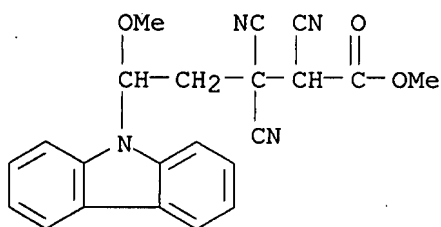
RN 102852-12-0 CAPLUS

CN 9H-Carbazole-9-butanoic acid, α -cyano- α -(dicyanomethyl)- γ -methoxy-, methyl ester (9CI) (CA INDEX NAME)



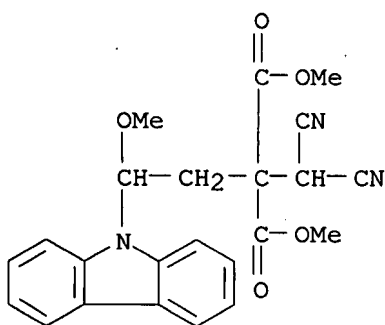
RN 102852-13-1 CAPLUS

CN 9H-Carbazole-9-pentanoic acid, α,β,β -tricyano- δ -methoxy-, methyl ester (9CI) (CA INDEX NAME)



RN 102852-14-2 CAPLUS

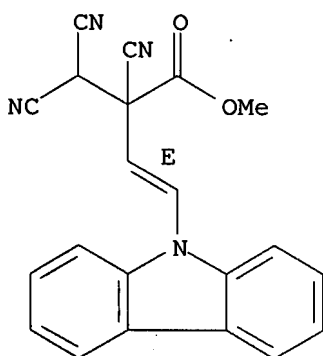
CN Propanedioic acid, [2-(9H-carbazol-9-yl)-2-methoxyethyl] (dicyanomethyl)-, dimethyl ester (9CI) (CA INDEX NAME)



RN 102852-38-0 CAPLUS

CN 3-Butenoic acid, 4-(9H-carbazol-9-yl)-2-cyano-2-(dicyanomethyl)-, methyl ester, (E)- (9CI) (CA INDEX NAME)

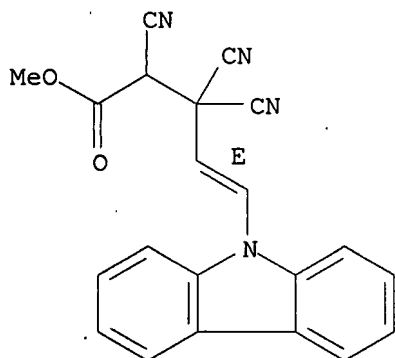
Double bond geometry as shown.



RN 102852-39-1 CAPLUS

CN 4-Pentenoic acid, 5-(9H-carbazol-9-yl)-2,3,3-tricyano-, methyl ester, (E)- (9CI) (CA INDEX NAME)

Double bond geometry as shown.



L6 ANSWER 55 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1985:406747 CAPLUS

DOCUMENT NUMBER: 103:6747

TITLE: Zwitterionic tetramethylene intermediates: a new interpretation for "charge-transfer" initiation

AUTHOR(S): Hall, H. K., Jr.; Gotoh, T.

CORPORATE SOURCE: Dep. Chem., Univ. Arizona, Tucson, AZ, 85721, USA

SOURCE: Polymer Preprints (American Chemical Society, Division of Polymer Chemistry) (1985), 26(1), 34-5

CODEN: ACPPAY; ISSN: 0032-3934

DOCUMENT TYPE: Journal

LANGUAGE: English

AB Investigation of the initiation mechanism in polymerization of N-vinylcarbazole (I) [1484-13-5] in the presence of tetracyanoethylene [670-54-2] or di-Me 2,2-dicyanoethylene-1,1-dicarboxylate [82849-49-8] showed that neither the I-cyano compound charge transfer complexes nor the ion-radical pairs formed from them initiated polymerization. The initiating species was the gauche or trans tetramethylene zwitterion formed as an intermediate from the charge-transfer complex. This finding indicated that cyclobutanes initiated vinyl polymerization. The mechanism and the kinetics of the zwitterionic initiation were discussed.

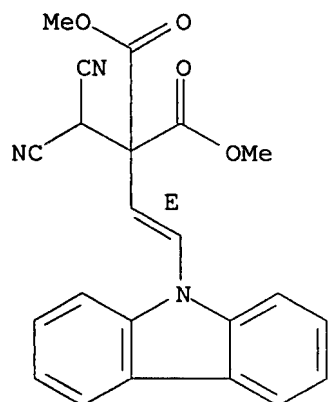
IT 96735-90-9

RL: CAT (Catalyst use); USES (Uses)
(catalysts, for vinylcarbazole polymerization)

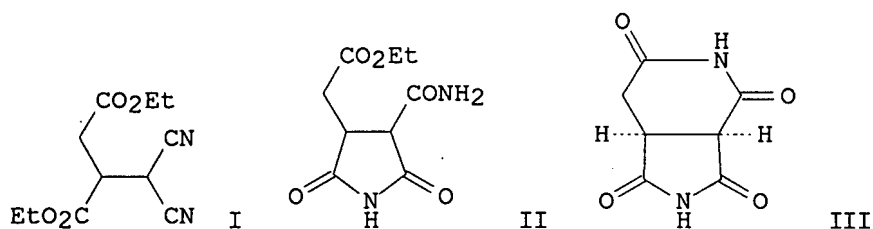
RN 96735-90-9 CAPLUS

CN Propanedioic acid, [2-(9H-carbazol-9-yl)ethenyl] (dicyanomethyl)-, dimethyl ester, (E)- (9CI) (CA INDEX NAME)

Double bond geometry as shown.



L6 ANSWER 56 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN
 ACCESSION NUMBER: 1983:488082 CAPLUS
 DOCUMENT NUMBER: 99:88082
 TITLE: Tetraoxo derivatives of perhydropyrrolo[3,4-c]pyridine
 AUTHOR(S): Ferappi, M.; Carotti, A.; Casini, G.; De Laurentis, N.; Giardina, D.; Cingolani, G. M.; Gavuzzo, E.; Mazza, F.
 CORPORATE SOURCE: Ist. Chim. Farm. Tossicol., Univ. Bari, Bari, 70126, Italy
 SOURCE: Journal of Heterocyclic Chemistry (1983), 20(2), 439-46
 CODEN: JHTCAD; ISSN: 0022-152X
 DOCUMENT TYPE: Journal
 LANGUAGE: English
 OTHER SOURCE(S): CASREACT 99:88082
 GI



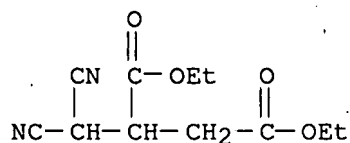
AB Michael adducts from di-Et furmarate with malonic esters or nitriles were cyclized to succinimide intermediates which, after glutarimide ring closure, afforded several N-Me and N-benzyl derivs. of cis-1,3,4,6-tetraoxoperhydropyrrolo[3,4-c]pyridine whose configuration was demonstrated by x-ray crystal structure anal. Thus, treating the adduct I with H₂SO₄ gave succinimide II which was treated with NaOEt in EtOH or tosyl acid in xylene to give pyrrolopyridine III.

IT 82584-86-9P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
 (preparation and cyclization of, pyrrolidine from)

RN 82584-86-9 CAPLUS

CN Butanedioic acid, (dicyanomethyl)-, diethyl ester (9CI) (CA INDEX NAME)



L6 ANSWER 57 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN
 ACCESSION NUMBER: 1982:582893 CAPLUS
 DOCUMENT NUMBER: 97:182893
 TITLE: Dimethyl 1,1-dicyanoethene-2,2-dicarboxylate, a new electrophilic olefin
 AUTHOR(S): Hall, H. K., Jr.; Sentman, R. C.
 CORPORATE SOURCE: Dep. Chem., Univ. Arizona, Tucson, AZ, 85721, USA
 SOURCE: Journal of Organic Chemistry (1982), 47(23), 4572-7
 CODEN: JOCEAH; ISSN: 0022-3263

DOCUMENT TYPE: Journal
LANGUAGE: English

AB dimethyl 1,1-dicyanoethene-2,2-dicarboxylate (I) [82849-49-8] was synthesized via a Knoevenagel condensation. I spontaneously copolymerizes with electron-rich olefins such as styrene [100-42-5] and p-methylstyrene [622-97-9]. In the copolymer, the bulky growing styryl radicals add to the dicyano-bearing carbon of I. Cyclobutane adducts are obtained in thermal reactions with styrene, p-methylstyrene, p-methoxystyrene [637-69-4], and vinyl ethers via a tetramethylene intermediate. Bond formation occurs at the diester end of I due to the greater stabilization provided by the dicyano group and the minimal steric requirements of the attacking methylene.

IT 82849-58-9P

RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation) (preparation and NMR spectra of)

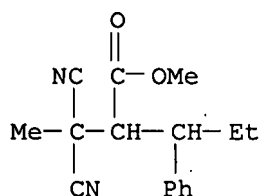
RN 82849-58-9 CAPLUS

CN Benzenepropanoic acid, α -(1,1-dicyanoethyl)- β -ethyl-, methyl ester, polymer with ethenylbenzene (9CI) (CA INDEX NAME)

CM 1

CRN 82917-40-6

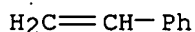
CMF C16 H18 N2 O2



CM 2

CRN 100-42-5

CMF C8 H8



L6 ANSWER 58 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1982:472220 CAPLUS

DOCUMENT NUMBER: 97:72220

TITLE: Contribution to the synthesis of the glutarimides.
III

AUTHOR(S): Victory, Pedro; Jover, Jose Maria; Sempere, Julian

CORPORATE SOURCE: Dep. Quim. Org., Inst. Quim. Sarria, Barcelona, Spain

SOURCE: Afinidad (1981), 38(376), 491-5

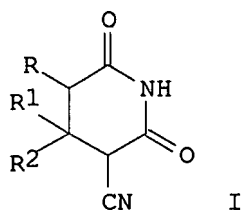
CODEN: AFINAE; ISSN: 0001-9704

DOCUMENT TYPE: Journal

LANGUAGE: Spanish

OTHER SOURCE(S): CASREACT 97:72220

GI

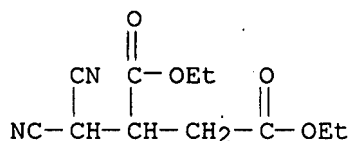


AB Glutarimides I [R = R1 = H, R2 = CO2Et, Ph, 3-furyl, 2-thienyl, Me; R = cyano, R1R2 = (CH2)5; R = Me, R1 = R2 = H] were prepared by treating CH2(CN)2 with R1R2C:CRCO2Et with or without isolation of (NC)2CHCR1R2CHRCO2Et, and acid hydrolysis of the enol ethers. Alternatively R1R2C:CRCO2Et was cyclized with NCCH2CONH2.

IT 82584-86-9P
 RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
 (preparation and cyclization of)

RN 82584-86-9 CAPLUS

CN Butanedioic acid, (dicyanomethyl)-, diethyl ester (9CI) (CA INDEX NAME)



L6 ANSWER 59 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1981:442180 CAPLUS

DOCUMENT NUMBER: 95:42180

TITLE: Absolute configuration of 2,7-diazaspiro[4,4]nonane. A reassignment

AUTHOR(S): Overberger, C. G.; Wang, David Wei; Hill, Richard K.; Krow, Grant R.; Ladner, David W.

CORPORATE SOURCE: Macromol. Res. Cent., Univ. Michigan, Ann Arbor, MI, 48109, USA

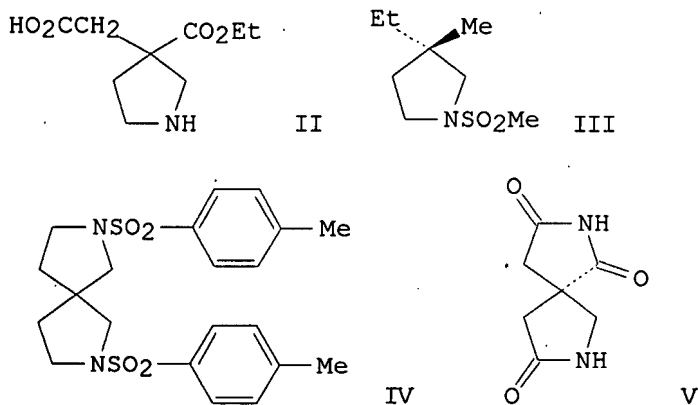
SOURCE: Journal of Organic Chemistry (1981), 46(13), 2757-64
 CODEN: JOCEAH; ISSN: 0022-3263

DOCUMENT TYPE: Journal

LANGUAGE: English

OTHER SOURCE(S): CASREACT 95:42180

GI



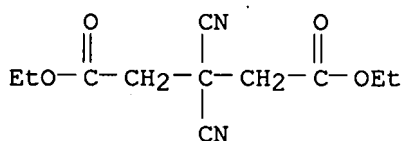
AB The absolute configuration of the axially dissym. spirane 2,7-diazaspiro[4,4]nonane (I), was elucidated as (R)-(-), (S)-(+) in CHCl₃ by synthesis of both enantiomers from the centrodissym. intermediate II; the configuration of (R)-(-)-II was correlated with that of (S)-HO₂CCMeEtCH₂CO₂H through the substituted pyrrolidine III. The configuration thus established for the sulfonamide derivative IV is opposite to that derived earlier (Krow, G. and Hill, R. K., 1968). The source of the original error lies in the preparation of spiroimide V, which is accompanied by almost total racemization when carried out at high temps. A more direct, efficient synthesis of I is described, followed by resolution with dinitrodiphenic acid to give the optically pure enantiomers. Lowe's rule predicts correctly the absolute configurations of several I derivs. but not that of I itself.

IT 77415-69-1P

RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation and diazaspirononane derivative from)

RN 77415-69-1 CAPLUS

CN Pentanedioic acid, 3,3-dicyano-, diethyl ester (9CI) (CA INDEX NAME)



L6 ANSWER 60 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1978:50444 CAPLUS

DOCUMENT NUMBER: 88:50444

TITLE: The chemistry of 2-oxopropanedinitrile (carbonyl cyanide); XIX. The ene synthesis using 2-oxopropanedinitrile and 1,3-dicarbonyl compounds

AUTHOR(S): Kociolek, K.; Leplawy, M. T.

CORPORATE SOURCE: Inst. Org. Chem., Tech. Univ. Lodz, Lodz, Pol.

SOURCE: Synthesis (1977), (11), 778-80

CODEN: SYNTBF; ISSN: 0039-7881

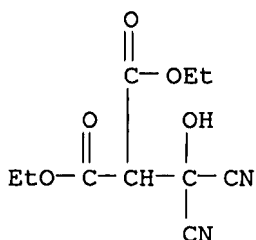
DOCUMENT TYPE: Journal

LANGUAGE: English

OTHER SOURCE(S): CASREACT 88:50444

AB Reaction of CO(CN)₂ with RCOCH₂COR₁ (I; R = R₁ = Ph, 2,4,6-Cl₃C₆H₂, Me; R = Me, F₃C, R₁ = Ph) in ether at 0° was complete in 1 h and gave RCOCH(COR₁)C(CN)₂OH (II; R and R₁ as before) in 100% yield. Reaction of CO(CN)₂ with I (R = R₁ = OEt) at room temperature required 20 days and gave II

in 43-66% yield.
 IT 65305-78-4P
 RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
 (Reactant or reagent)
 (preparation and reaction of, with aniline)
 RN 65305-78-4 CAPLUS
 CN Propanedioic acid, (dicyanohydroxymethyl)-, diethyl ester (9CI) (CA INDEX
 NAME)

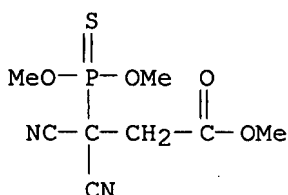


L6 ANSWER 61 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN
 ACCESSION NUMBER: 1973:545978 CAPLUS
 DOCUMENT NUMBER: 79:145978
 TITLE: O,O-Dialkylthiophosphoric acid pseudochalcogen acyls
 INVENTOR(S): Koehler, Helmut; Gerats, Irmtraut; Eichler, Gerhard;
 Kochmann, Werner
 SOURCE: Ger. (East), 14 pp.
 CODEN: GEXXA8
 DOCUMENT TYPE: Patent
 LANGUAGE: German
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
DD 95374	A1	19730212	DD 1971-156303	19710705 <--
PRIORITY APPLN. INFO.:			DD 1971-156303	A1 19710705

AB (MeO)2P(S)N(CN)CH2CO2R (I) and/or (MeO)2P(:NCN)SCH2CO2R (II) (R = Me or Et), prepared by reacting (MeO)2P(S)NNaCN with XCH2CO2R (X = Br or Cl), gave 95.0, 52.5 and 69.0% mortality for R = Me and 92.5, 51.0 and 55.0% for R = Et at 0.01, 1.0 and 0.05 weight % concentration, resp., against *Musca domestica*, *Sitophilus granarius* and *Tetranychus urticae*, resp. Analogs of I and II wherein the CO2R group was replaced by CONH2 and CONHMe, and (MeO)2P(S)C(CN)2CH2COR and (MeO)2P[:C(CN)2]SCH2COR (R = NHMe or OMe) were also prepared

IT 50605-40-8P
 RL: SPN (Synthetic preparation); PREP (Preparation)
 (preparation of)
 RN 50605-40-8 CAPLUS
 CN Propanoic acid, 3,3-dicyano-3-(dimethoxyphosphinothioyl)-, methyl ester (9CI) (CA INDEX NAME)



L6 ANSWER 62 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1973:545936 CAPLUS

DOCUMENT NUMBER: 79:145936

TITLE: Reaction of some fluoroolefins with sodium cyanide

AUTHOR(S): Dyatkin, B. L.; Sterlin, S. R.; Zhuravkova, L. G.;
Martynov, B. I.; Knunyants, I. L.

CORPORATE SOURCE: Inst. Elementoorg. Soedin., Moscow, USSR

SOURCE: Zhurnal Organicheskoi Khimii (1973), 9(9),
1786-90

CODEN: ZORKAE; ISSN: 0514-7492

DOCUMENT TYPE: Journal

LANGUAGE: Russian

GI For diagram(s), see printed CA Issue.

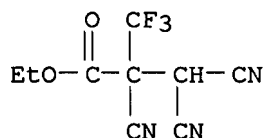
AB (CF₃)₂C:CF₂ reacted with 1 equiv of NaCN at -5 to 0° in dioxane containing H₂O to give 38% (CF₃)₂C:CFCN (I), and with excess NaCN in THF containing H₂O to give 55% NCC(CF₃)₂CH(CN)₂, a C-H acid of pK_a 2.12; analogous treatment of (CF₃)₂C:CFPh and (CF₃)₂CHCO₂Et yielded 49% NCC(CF₃)₂CHPhCN and 51% EtO₂CC(CN)(CF₃)CH(CN)₂, resp., after neutralization. Under similar conditions, CF₃CF:CF₂ afforded 59% CF₃[C(CN)₂]₂Na, although its acid could not be isolated, and (CF₃)₂C:CFOEt gave 3% (CF₃)₂C:C(CN)OEt. I reacted with H₂SO₄ and EtOH to give 25% (CF₃)₂C:CFCO₂Et, with HCl in EtOH to give 20% HOC(CF₃)₂CHFCONH₂, with Et₂NH to give 43% (CF₃)₂C:C(CN)NEt₂, with PhNH₂ to give 60% (CF₃)₂CHC(CN):NPh, and with concentrated H₂SO₄ to give 84% iminolactone (II; R = H), which was converted to its Hg salt (II; R = 1/2 Hg) with HgO in refluxing aqueous Me₂CO.

IT 50616-04-1P

RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of)

RN 50616-04-1 CAPLUS

CN Propanoic acid, 2-cyano-2-(dicyanomethyl)-3,3,3-trifluoro-, ethyl ester
(9CI) (CA INDEX NAME)



L6 ANSWER 63 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1965:462918 CAPLUS

DOCUMENT NUMBER: 63:62918

ORIGINAL REFERENCE NO.: 63:11492h

TITLE: Reaction of acetylenic esters with cyanoacetic ester
and pyridine

AUTHOR(S): Bamfield, P.; Crabtree, A.; Johnson, A. W.

CORPORATE SOURCE: Univ. Nottingham, UK

SOURCE: Journal of the Chemical Society (1965)
4355-62

CODEN: JCSOA9; ISSN: 0368-1769

DOCUMENT TYPE: Journal

LANGUAGE: English

AB Modified structures are suggested for the yellow and the blue adducts from dimethyl acetylenedicarboxylate, Et cyanoacetate, and pyridine, which were originally prepared and formulated by Diels. The reaction of Me phenylpropiolate, Et cyanoacetate, and pyridine leads to a 1:1:1-adduct in which the pyridine has suffered ring-fission. Various reactions of the adducts are discussed.

IT 1289-25-4

(Derived from data in the 7th Collective Formula Index (1962-1966))

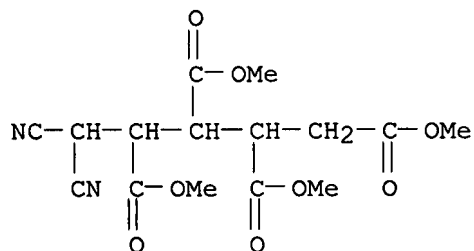
RN 1289-25-4 CAPLUS

CN 1,2,3,4-Pentanetetracarboxylic acid, 5,5-dicyano-, tetramethyl ester, compd. with pyridine (1:1) (8CI) (CA INDEX NAME)

CM 1

CRN 45287-28-3

CMF C15 H18 N2 O8



CM 2

CRN 110-86-1

CMF C5 H5 N



L6 ANSWER 64 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1964:484793 CAPLUS

DOCUMENT NUMBER: 61:84793

ORIGINAL REFERENCE NO.: 61:14826g-h,14827a-c

TITLE: 1-Halo-1,2,3,3-tetra(negatively substituted)propanes and their salts

INVENTOR(S): Martin, Elmore L.

PATENT ASSIGNEE(S): E. I. du Pont de Nemours & Co.

SOURCE: 6 pp.

DOCUMENT TYPE: Patent

LANGUAGE: Unavailable

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 3133084		19640512	US	19600624 <--
PRIORITY APPLN. INFO.:			US	19600624

AB Compds. of the general formula $[XC(Z):C(Y)C(A)R]-M^+$ (I), where A, R, Y, Z are electron withdrawing groups such as CN, CO₂Et, Bz, or SO₂Ph, X is Cl or F, and M is H, Na, K, or a substituted ammonium ion, are dyes for natural and synthetic fibers. Thus, H₂C(CN)₂ 79 in tetrahydrofuran (II) 220 was added with stirring to a dispersion 52 of 51.2% NaH in mineral oil and II 660 at 5-10° during 15 min., the mixture stirred 30 min., then dichlorofumaronitrile 44 in II 220 added during 15 min., II vacuum-distilled at 35-40°, the residual yellow solid dissolved in H₂O 250, the pH adjusted to 8 with CO₂, then Et₄NBr 100 in H₂O 200 parts added slowly with

stirring, the mixture cooled to 5°, and the yellow crystals of I (A = R = Y = X = CN, Z = Cl, M = Et4N) (III) filtered, washed with 1% Et4NBr, and then H2O. The cake was dissolved in H2O 3500 at 100°, decolorizing carbon 10 added, the solution clarified, cooled to 5°, the long yellow needles filtered, washed with H2O and air-dried, giving 70 parts III, m. 129-31°, λ_{maximum} 387 mμ, ε = 18,200

(MeOH) yellow on cellulose acetate and nylon, brownish yellow on wool and silk. Similarly, other I were prepared as tabulated below: X, Z, Y, A, R, M, % yield, m.p., color, λ (mμ)maximum, ε; Cl, PhN(CO-)2, CN, CN, Me4N, 31 230-5° (decompose), orange, 468, 12,200; Cl, CO2Me, CO2Me, CN, CN, Et4N, 82, 88-90°, yellow, 335, 29,400; Cl, CN, CN, CO2Et, CO2Et, H, 100, bl, 115-20°, yellow (Na salt), -, -; Cl, Bz, Bz, CN, CN, Me4N, 39, 210-12° (decompose), yellow, 416, 27,000; F, CF3, CF3, CN, CN, Pr4N, 81, 84-6°, yellow, -, -; Cl, CN, CN, CN, CN, Me4N, -, 217-18° (decompose), yellow, 386, 17,600; Cl, CN, CN, CN, CN, Pr4N, -, 74-6° (decompose), yellow, 386, 18,100; Cl, CN, CN, CN, CN, Et3NH, -, 63-5° (decompose), yellow, 387, 17,200; Cl, CN, CN, CN, CO2Et, Et4N, 56, 70-2°, yellow, 400, 15,700; Cl, CN, CN, CN, SO2C6H4Me-4, Me4N, 73, 124-6° (decompose), yellow, 387, 17,000; Cl, CN, CN, CN, Bz, Me4N, -, 159-61°, yellow, 414, 17,100; Cl, CN, CN, CN, Bz, Et4N, 30 118-19°, yellow, 420, 16,200; Cl, CN, CN, CN, CN, Pr4N, -, 109-10°, yellow, 412, 17,600; Cl, CF3, CF3, CN, CN, Et4N, 64, 84-5°, yellow, -, -; F, -CF2CF2-, CN, CN, Na, -, -, orange, -, -; Cl, CN, CN, Bz, Bz, Me4n, -, 167°9°, yellow, 422, 8000; Cl, CN, CN, CN, CONHPh, K, -, -, red, -, -; Cl, CN, CN, SO2Ph, SO2Ph, Me4N, -, -, yellow, -, -; Cl, CN, CN, Bz, CO2Et, H, 20, 97-8°, colorless, -, -; Cl, CN, CN, Bz, CO2Et, Na, yellow;

IT 98469-37-5P, Ammonium, tetraethyl, 1,2-dicarboxy-1-chloro-3,3-dicyanopropenide, dimethyl ester

RL: PREP (Preparation)

(preparation of)

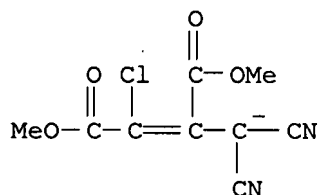
RN 98469-37-5 CAPLUS

CN Tetraethylammonium 1,2-dicarboxy-1-chloro-3,3-dicyanopropenide, dimethyl ester (7CI) (CA INDEX NAME)

CM 1

CRN 98469-36-4

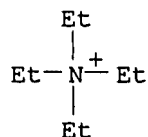
CMF C9 H6 Cl N2 O4



CM 2

CRN 66-40-0

CMF C8 H20 N



L6 ANSWER 65 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1962:429396 CAPLUS

DOCUMENT NUMBER: 57:29396

ORIGINAL REFERENCE NO.: 57:5809h-i,5810c

TITLE: Nitration of cyclohexanecarboxylic acid to caprolactam

AUTHOR(S): Bigot, J. A.; Meijerink, Th. A. J.; Revallier, L. J.

CORPORATE SOURCE: Central Lab., Staatsmijnen, Geleen, Neth.

SOURCE: Recueil des Travaux Chimiques des Pays-Bas (1962), 81, 363-4

CODEN: RTCPA3; ISSN: 0165-0513

DOCUMENT TYPE: Journal

LANGUAGE: English

AB Cyclohexanecarboxylic acid (I) with nitryl hydroxyrosulfate in oleum did not give the expected nitrocyclohexane, but 70% caprolactam (II) and a mixture of m-dinitrobenzene and nitrobenzene (total yield 22%, based on I). The mechanism of the reaction is unknown, but there is some evidence that removal of H₂O from a nitro derivative is 1 of the steps involved. 1-Methyl-1-nitrocyclohexane with oleum gave a compound, C₇H₁₁NO (b₂ 79°, m. 48°), probably 1-methyl-1-nitrocyclohexene (or its rimer), a compound that could be isolated as such, since it could neither dehydrogenate to a C₆H₆ derivative, nor disproportionate and subsequently rearrange to II.

IT 94211-18-4P, Acrylic acid, 3,3-dicyano-2-(dicyanomethyl)-, ethyl ester, compound with quinoline 94467-89-7P, Acrylic acid, 3,3-dicyano-2-(dicyanomethyl)-, ethyl ester, compound with NH₃
RL: PREP (Preparation)
(preparation of)

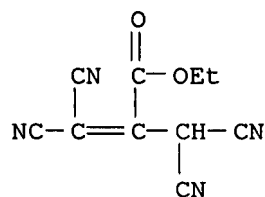
RN 94211-18-4 CAPLUS

CN Acrylic acid, 3,3-dicyano-2-(dicyanomethyl)-, ethyl ester, compd. with quinoline (7CI) (CA INDEX NAME)

CM 1

CRN 94211-17-3

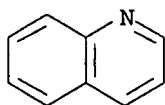
CMF C10 H6 N4 O2



CM 2

CRN 91-22-5

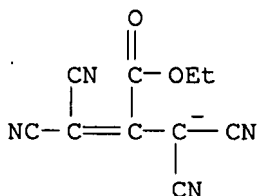
CMF C9 H7 N



RN 94467-89-7 CAPLUS

CN 2-Propenoic acid, 3,3-dicyano-2-(dicyanomethyl)-, ethyl ester, ion(1-),

ammonium (9CI) (CA INDEX NAME)



● NH₄⁺

L6 ANSWER 66 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1962:429395 CAPLUS

DOCUMENT NUMBER: 57:29395

ORIGINAL REFERENCE NO.: 57:5809e-h

TITLE: Base-catalyzed ring opening of diethyl
1,1,2,2-tetracyanocyclopropane-3,3-dicarboxylate
Regan, T. H.

AUTHOR(S): E. I. du Pont de Nemours & Co., Wilmington, DE

CORPORATE SOURCE: Journal of Organic Chemistry (1962), 27,
2236-7

SOURCE: CODEN: JOCEAH; ISSN: 0022-3263

DOCUMENT TYPE: Journal

LANGUAGE: Unavailable

AB An example of a cyclopropane ring cleavage under very mild conditions was reported. CH₂(CN)₂ (6.6 g.) in 17.4 g. di-Et oxomalonate was left 3 hrs. with one drop of base catalyst; the solid was collected and shown to be di-Et dihydroxymalonate. Fractionation of the yellow oil gave 11.7 g. di-Et 1,1-dicyanoethylene-2,2-dicarboxylate, b₁ 86°, n_D 1.4628. An equimolar mixture of this compound and anthracene after heating at 150° gave crystalline product, m. 153.6-5.2° (alc.-H₂O). The product resulting from 38 g. CH₂(CN)₂ and 100 g. di-Et oxomalonate in 250 ml. alc. treated in the cold with 52 g. Br, the solution poured onto 1 kg. ice, and the oil crystallized when left overnight gave 71.5 g. di-Et 1,1,2,2-tetracyanocyclopropane-3,3-dicarboxylate (I), m. 129.6-31.2° (alc.-H₂O). I (15 g.) was suspended in 500 ml. Et₂O, treated with 15 g. dry NH₃, stirred overnight and the mixture filtered to give 11.4 g. solid, m. 192-201° (decomposition). The filtrate evaporated and the residue stirred with CHCl₃ gave 0.5 g. yellow powder, m. 203° (decomposition). The CHCl₃ solution evaporated gave Et carbamate, m. 46.6-8.6°. The yellow powder was ammonium 1,1,3,3-tetracyano-2-carbethoxypropenide (II). II in H₂O treated with a concentrated aqueous solution of

quinolinium chloride gave quinolinium 1,1,3,3-tetracyano-2-carbethoxypropenide, m. 111.5-12.5°. Recrystn. from H₂O gave a hydrate, m. 51-2°.

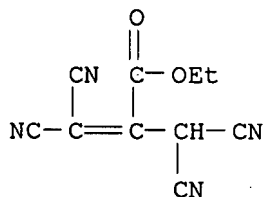
IT 94211-18-4P, Quinoline, compound with Et 3,3-dicyano-2-(dicyanomethyl)-acrylate 94467-89-7P, Acrylic acid, 3,3-dicyano-2-(dicyanomethyl)-, ethyl ester, compound with NH₃
RL: PREP (Preparation)
(preparation of)

RN 94211-18-4 CAPLUS

CN Acrylic acid, 3,3-dicyano-2-(dicyanomethyl)-, ethyl ester, compd. with quinoline (7CI) (CA INDEX NAME)

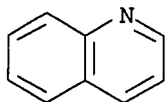
CM 1

CRN 94211-17-3
CMF C10 H6 N4 O2

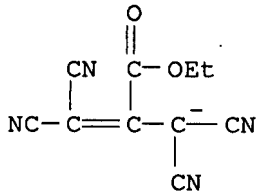


CM 2

CRN 91-22-5
CMF C9 H7 N



RN 94467-89-7 CAPLUS
CN 2-Propenoic acid, 3,3-dicyano-2-(dicyanomethyl)-, ethyl ester, ion(1-), ammonium (9CI) (CA INDEX NAME)



● NH₄⁺

L6 ANSWER 67 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN
ACCESSION NUMBER: 1940:18276 CAPLUS
DOCUMENT NUMBER: 34:18276
ORIGINAL REFERENCE NO.: 34:2801g-i,2802a-b
TITLE: Synthesis of α,α -dimethyltricarballic and 1-carboxy-cyclopentane-1- α -succinic and 1-carboxy-3-methylcyclopentane-1- α -succinic acids
AUTHOR(S): Desai, R. D.; Sahariya, G. S.
SOURCE: Journal of the University of Bombay, Science: Physical Sciences, Mathematics, Biological Sciences and Medicine (1939), 8(Pt. 3), 235-8
CODEN: JUBSAS; ISSN: 0368-4644
DOCUMENT TYPE: Journal
LANGUAGE: Unavailable
AB When in RR'C(CN)C(CN)CO₂Et, R and R' together are cyclopentane or

methylocyclopentane rings, there are obtained with CH₂BrCO₂Et (I) excellent yields of tricarballic acids which are characterized by their toluidide N-tolylimides. A mixture of the Et sodiocyanoacetate (II) from 24 g. cyanoacetate and 21 g. cyclopentanone cyanohydrin is allowed to stand for 48 h. After addition of 32 g. I the mixture is kept at room temperature for 2 days

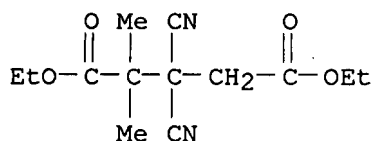
and then refluxed until it is neutral. The EtOH is distilled off, the residue diluted with H₂O and the oil extracted with ether, dried and distilled into

3 fractions, b₄ 90-120°, 120-65° and 185-7°. The 2nd fraction is retreated with I. Et 1-cyanocyclopentane-1-α-cyanosuccinate (III), b₄ 185-7°, is obtained in 45% yield. Hydrolysis of III with concentrated H₂SO₄ gives 1-carboxycyclopentanesuccinic acid (IV), m. 165° (cf. Chatterji, C. A. 31, 7409.7, found 159°). Its anilide N-phenylimide, prepared by heating IV with PhNH₂ at 170-5° for 3 h., m. 156°; p-toluidide N-p-tolylimide m. 189-90°. Et 1-cyano-3-methylocyclopentane-1-α-cyanosuccinate (V), prepared as II, b₁₂ 205°. Hydrolysis of V gives 1-carboxy-3-methylocyclopentanesuccinic acid, m. 144°; its p-toluidide N-tolylimide m. 167°. Di-Et 2-methyl-3,3-dicyanobutane-3,4-dicarboxylate (VI), prepared from II, Me₂C(OH)CN and I, in 45% yield, b₅ 176-8°. Saponification of VI with H₂SO₄ gives α,α-dimethyltricarballic acid, m. 160° (C. found 156°). Its anilide N-phenylimide m. 140°; the p-toluidide N-p-tolylimide m. 170°.

IT 858794-64-6P, Glutaric acid, β,β-dicyano-α,α-dimethyl-, diethyl ester
RL: PREP (Preparation)
(preparation of)

RN 858794-64-6 CAPLUS

CN Glutaric acid, β,β-dicyano-α,α-dimethyl-, diethyl ester (4CI) (CA INDEX NAME)



L6 ANSWER 68 OF 68 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1938:911 CAPLUS

DOCUMENT NUMBER: 32:911

ORIGINAL REFERENCE NO.: 32:156d-i,157a-i,158a-e

TITLE: 2,3,-Dioxopyrrolines, mononuclear substances related to isatin

AUTHOR(S): Mumm, Otto; Hornhardt, Hans

SOURCE: Berichte der Deutschen Chemischen Gesellschaft
[Abteilung] B: Abhandlungen (1937), 70B,
1930-47

CODEN: BDCBAD; ISSN: 0365-9488

DOCUMENT TYPE: Journal

LANGUAGE: Unavailable

AB 5-Phenyl-2,3-dioxopyrroline (I) (C. A. 5, 703) is so extraordinarily similar in appearance and chemical properties to isatin that it may be considered as a mononuclear isatin, and it was hoped that by means of this very reactive substance light might be obtained on some of the controversial questions on isatin, especially the structure of its salts and derivs. The earlier work on I was accordingly resumed and attention was directed to the preparation of analogs of I containing an aliphatic residue instead of

Ph. In order not to weaken the ring unnecessarily, a residue (hexyl) of rather high mol. weight was chosen. As a tertiary residue might also favor the stability of the ring, Me₃C was selected for a 2nd series of expts. It was intended to prepare the new compds. by the earlier method. Pinacolone and C₆H₁₃COMe were condensed with HCO₂R to the hydroxymethylene compds. which with NH₂OH yielded the oxazoles through the intermediate oximes. The conversion of the oxazoles into the open-chain methylimide nitriles proceeded as expected and the formation from the nitriles of the desired pyrrolines with alc. HCl undoubtedly occurred, as evidenced by the appearance of the characteristic dark red color, but the products did not crystallize. The planned investigation was therefore continued with aromatic derivs., using p-tolyl instead of Ph compds. These tolyl compds., having higher m. ps., were considerably more stable and crystallized better. The starting point was α-p-tolylisoxazole (II). That the nitrile obtained from II was really p-tolylpyruvonnitrile methylimide, RC(OH):CHC(:NMe)CN(R = p-MeC₆H₄) (III), was shown by the reaction with MeMgI, which gave the normal product, RC(OH)MeCH₂C(:NMe)C(:NMgI) (IV), and also the compound RC(NH₂)MeCH₂C(:NMe)C(:NMgI)Me(V) when the Grignard compound was decomposed with NH₄Cl instead of water. RC(OH)MeCH₂C(:NMe)C(OH)(NHMgI)Me (VI) was also formed by addition of H₂O to IV under the influence of glacial AcOH. III merely treated in the cold with HCl in absolute alc. gave the blood-red di-HCl salt of 5-p-tolyl-2-oxo-3-methyliminopyrroline (VII). The previously assumed intermediate imido ester, RC(OH):CHC(:NMe)C(:NH)OEt (VIII), corresponding to the nitrile, was isolated as its white HCl salt, which readily changes, even in the absence of air, into the dark red derivative of VII. The distribution of the double bonds shown in III probably occurs only under the influence of the HCl, the free nitrile having the tautomeric structure RCOCH:C(NHMe)CN. The outstanding property of the dark red VII.2HCl is the ease with which the NMe group is replaced by O to form the brick-red 5-p-tolyl-2,3-dioxopyrroline (IX). As with isatin, NaOH cleaves the ring in IX to give α-oxo-γ-imino-γ-p-tolylbutyric acid (X) through an intermediate intensely blue alkali salt. Attempts to liberate VII from its HCl salt were unsuccessful. Dilute aqueous alkali or NaHCO₃ gave, instead, the yellow-green pseudo base (XI), while excess of concentrated KOH yielded a dark red K salt, Cl₂H₁₁ON₂K.2H₂O, which regenerated XI with water. NH₃ in alc. replaces both the NMe group and the carbonyl O by NH and at the same time 1 mol. alc. is taken up with formation of a product, RC:CH.C(NH₂)(OEt).C(:NH).NH (XII), similar in structure to XI; the dark red color immediately produced by HCl shows the ring has not been cleaved. PhNH₂ in alc. yields brick-red needles of the 3-phenylimino analog (XIII) of VII. With KOH and also with HCl, XIII forms salts which are red-violet in solution and almost black in the solid state. The HCl salt quant. splits off the HCl at high temps. in vacuo without changing to the brick-red of the free XIII, showing that the salt formation is accompanied by a simultaneous intramol. rearrangement. In water the HCl salt, like that of VII, is hydrolyzed to IX, but attempts to prepare the pseudo base were unsuccessful; instead was obtained XIII into which the K salt also changes on mere exposure to moist air. This difference in behavior and the very different colors show that the salts of VII and XIII have different structures. As with water and PhNH₂, the dark red salt of VII also reacts with compds. having a reactive methylene group. Especially smooth, and under the mildest conditions, is the reaction with CH₂(CN)₂ to give 5-p-tolyl-2-oxo-3-dicyanomethylenepyrroline (XIV), also obtained from IX or XIII. Surprisingly, XIV forms beautiful violet-black needles and dissolves, although difficultly, in alc. with red-violet color, whereas the corresponding isatin derivative is yellow-red, indicating a fundamental difference in structure. When the alc. solution of XIV is treated with a strong base, it immediately turns steel-blue, but, as with the salts of IX, the blue color quickly disappears and the ring is opened; acids precipitate the yellow cleavage product,

RC(NH₂):CHC[:C(CN)₂]CO₂H,
m. 276°, probably in the form of the inner salt, which with boiling

alc. HCl changes through the intermediate $RC(NH_2):CHC(CO_2H):C(CO_2H)C(:NH)OEt$ into the compound $RC(NH_2):CHC(CO_2H):CHC(:NH)OEt$ (XV). X gently heated with dilute acids yields the dioxo acid. The ring in IX is also cleaved by piperidine, $MeNH_2$ and NH_3 to form compds. of the type $RC(:NH)CH_2COCONC_5H_{11}$ (XVI). $PhNH_2$ and $CH_2(CN)_2$, on the other hand, react with the 3-CO group, leaving the ring intact. The similarity of the dioxopyrrolines to isatin is also shown in their catalytic hydrogenation. There is first formed a light gray, alc.-insol. product (XVII) corresponding to isatyde which in the air rapidly regenerates the original compound. If the hydrogenation is continued, the XVII redissolves, and cautious addition of water to the colorless alc. solution ppts. a completely air-stable crystalline product, $RC(NH_2):CHCH(OH)CO_2H$ (XVIII). Reduction of XIII in alc. proceeds 1 step further, with addition of 2 mols. H and 1 mol. alc. to give the compound $RCH(NH_2)CH_2CH(NHPh)CO_2Et$ (XIX). For the bearing of the above facts, especially the color phenomena, on the structures of the mononuclear isatins and their derivs., the original should be consulted.

Hydroxymethylenepinacolone dioxime (54% yield), m. 84° .

α -tert-Butylisoxazole, b 760 156° ; its methosulfate with KCN in water at 0° gave 82% trimethylacetopyruvonnitrile methylimide, m. 42° , hydrolyzed by cold concentrated HCl to the pyruvic acid, crystals with 1 H_2O , m. 64° , and by dilute HCl to the amide, m. 115° ; in cold absolute alc. with HCl gas the nitrile imide gave a dark red oil which with 2 N NaOH or 50% AcOH yielded α -oxo- α -imino- δ,δ -dimethylcaproic acid, m. 185° (gas evolution).

When the red oil was carefully freed from adhering HCl, simple solution in ordinary alc. resulted in ring cleavage (probably by the water in the alc.), but AcOEt precipitated a crystalline substance, m. 186° , insol. in all solvents except alc. and water, which on gentle warming with water gave trimethylacetopyruvic acid methylimide, m. 183° .

Hydroxymethylenemethyl hexyl ketone oxime, m. 118° .

α -Hexylisoxazole, b 11 $97-8^\circ$, was analyzed as the chloroplatinate, $C_{20}H_{36}O_2N_2PtCl_6$, obtained from the methosulfate with $PtCl_4$. α,α -Dioxodecanonitrile α -methylimide, oil

decomposing on distillation, even in a high vacuum; α,α -dioxodecanamide, m. 99° . Hydroxymethylene-p-methylacetophenone oxime (67% yield), m. 133° . II, m. 60° . III, light yellow, m. 126° . IV (3 g. from 2 g. II and 2.2 mol. $MeMgI$ boiled 2 h. in ether), yellow, m. 175° (decomposition); heated a short time or allowed to stand 1 day at room temperature in glacial AcOH, it changed into VI, rhombic tables, red-brown in incident light, m. 183° (decomposition). V, m. 197° , soluble in AcOH with wine-red color, easily soluble in dilute HCl and repptd. by NaOH. VII.2HCl (78%), sinters and carbonizes at 183° ; picrate, intensely red, m. 192° . VIII.HCl, from III in cold

dioxane with 0.662 N HCl in absolute alc., decomp. 145° . If in the treatment of III with alc.-HCl water is present even only in traces the reaction proceeds in part in an entirely different way, giving in addition to the dark red salt Me p-toluylypyruvate, m. 84° ; free acid, crystals with 1 H_2O , m. 143° . Ag salt of VII, red needles with 1 MeOH, decomposing 172° . Cu salt, $(Cl_2H_{11}ON_2)_2Cu.4H_2O$, green, m. 191° (decomposition). XII, m. 153° . IX, precipitated quant. in about 6 h. from VII.2HCl in 20 parts cold water, m. $229-30^\circ$; a cold alc. suspension treated with somewhat less than 1 mol. EtOK-solution at once becomes

blue-violet and soon deposits the K salt, $Cl_{11}H_{80}N_2K.2H_2O$, which is not very stable even when dry; one sample had become yellowish after 14 days. Alc. IX treated with aqueous NaOH also immediately turns blue-violet but the color rapidly disappears and on cautious acidification X, m. 155° , seps. Piperidine (XVI), m. 184° . Amide, $Cl_{11}H_{120}N_2O.0.5H_2O$, m. 179° . Methylamide ($0.5H_2O$), m. 169° . XVIII, turns brown and carbonizes $245-50^\circ$. XIII, m. 237° . XIX, m. 123° . XIV was obtained in 92% yield; its melting or decomposition point is so extraordinarily high that it could not be determined XV.2HCl, lemon-yellow, m. $148-9^\circ$.

IT 855234-21-8P, 3-Butenoic acid, 4-amino-2-(dicyanomethylene)-4-p-
 tolyl-
 RL: PREP (Preparation)
 (preparation of)
 RN 855234-21-8 CAPLUS
 CN 3-Butenoic acid, 4-amino-2-(dicyanomethylene)-4-p-tolyl- (4CI) (CA INDEX
 NAME)

